47

(54) Title: ENDOTHELIN-1 PROMOTER POLYMORPHISM

(57) Abstract: Disclosed are single nucleotide polymorphisms (SNPs) associated with hypertension, end stage renal disease due to hypertension, non-insulin dependent diabetes mellitus, end stage renal disease due to non-insulin dependent diabetes mellitus, lung cancer, breast cancer, prostate cancer, colon cancer, atherosclerotic peripheral vascular disease due to hypertension, cerebrovascular accident due to hypertension, cataracts due to hypertension, cardiomyopathy with hypertension, myocardial infarction due to hypertension, atherosclerotic peripheral vascular disease due to non-insulin dependent diabetes mellitus, cerebrovascular accident due to non-insulin dependent diabetes mellitus, ischemic cardiomyopathy, ischemic cardiomyopathy with non-insulin dependent diabetes mellitus, myocardial infarction due to non-insulin dependent diabetes mellitus, atrial fibrillation without valvular disease, alcohol abuse, anxiety, asthma, chronic obstructive pulmonary disease, cholecystectomy, degenerative joint disease, end stage renal disease and frequent de-clots, end stage renal disease due to focal segmental glomerular sclerosis, end stage renal disease due to insulin dependent diabetes mellitus, or seizure disorder. Also disclosed are methods for using the SNPs to determine susceptibility to these diseases; nucleotide sequences containing the SNPs; kits for determining the presence of the SNPs; and methods of treatment or prophylaxis based on the presence of the SNPs.

ENDOTHELIN -1 PROMOTER POLYMORPHISM
BACKGROUND

This invention relates to detection of individuals at risk for pathological conditions based on the presence of single nucleotide polymorphisms (SNPs) at positions 2239 and 2657 on the human endothelin-1 (EDN-1) promoter.

During the course of evolution, spontaneous mutations appear in the genomes of organisms. It has been estimated that variations in genomic DNA sequences are created continuously at a rate of about 100 new single base changes per individual (Kondrashow, *J. Theor. Biol.*, 175:583-594, 1995; Crow, *Exp. Clin. Immunogenet.*, 12:121-128, 1995). These changes, in the progenitor nucleotide sequences, may confer an evolutionary advantage, in which case the frequency of the mutation will likely increase, an evolutionary disadvantage in which case the frequency of the mutation is likely to decrease, or the mutation will be neutral. In certain cases, the mutation may be lethal in which case the mutation is not passed on to the next generation and so is quickly eliminated from the population. In many cases, an equilibrium is established between the progenitor and mutant sequences so that both are present in the population. The presence of both forms of the sequence results in genetic variation or polymorphism. Over time, a significant number of mutations can accumulate within a population such that considerable polymorphism can exist between individuals within the population.

Numerous types of polymorphisms are known to exist. Polymorphisms can be created when DNA sequences are either inserted or deleted from the genome, for example, by viral insertion. Another source of sequence variation can be caused by the presence of repeated sequences in the genome variously termed short tandem repeats (STR), variable number tandem repeats (VNTR), short sequence repeats (SSR) or microsatellites. These repeats can be dinucleotide, trinucleotide, tetranucleotide or pentanucleotide repeats. Polymorphism results from variation in the number of repeated sequences found at a particular locus.

By far the most common source of variation in the genome is the single nucleotide polymorphism or SNP. SNPs account for approximately 90% of human DNA polymorphism (Collins et al., *Genome Res.*, 8:1229-1231, 1998). SNPs are single base pair positions in genomic DNA at which different sequence alternatives (alleles) exist in a population. In addition, the least frequent allele must occur at a frequency of 1% or greater. Several definitions of SNPs exist in the literature (Brooks, *Gene*, 234:177-186,

1999). As used herein, the term "single nucleotide polymorphism" or "SNP" includes all single base variants and so includes nucleotide insertions and deletions in addition to single nucleotide substitutions (e.g. A->G). Nucleotide substitutions are of two types. A transition is the replacement of one purine by another purine or one pyrimidine by another pyrimidine. A transversion is the replacement of a purine for a pyrimidine or vice versa.

The typical frequency at which SNPs are observed is about 1 per 1000 base pairs (Li and Sadler, *Genetics*, 129:513-523, 1991; Wang et al., *Science*, 280:1077-1082, 1998; Harding et al., *Am. J. Human Genet.*, 60:772-789, 1997; Taillon-Miller et al., *Genome Res.*, 8:748-754, 1998). The frequency of SNPs varies with the type and location of the change. In base substitutions, two-thirds of the substitutions involve the C<->T (G<->A) type. This variation in frequency is thought to be related to 5-methylcytosine deamination reactions that occur frequently, particularly at CpG dinucleotides. In regard to location, SNPs occur at a much higher frequency in non-coding regions than they do in coding regions.

SNPs can be associated with disease conditions in humans or animals. The association can be direct, as in the case of genetic diseases where the alteration in the genetic code caused by the SNP directly results in the disease condition. Examples of diseases in which single nucleotide polymorphisms result in disease conditions are sickle cell anemia and cystic fibrosis. The association can also be indirect, where the SNP does not directly cause the disease but alters the physiological environment such that there is an increased likelihood that the patient will develop the disease. SNPs can also be associated with disease conditions, but play no direct or indirect role in causing the disease. In this case, the SNP is located close to the defective gene, usually within 5 centimorgans, such that there is a strong association between the presence of the SNP and the disease state. Because of the high frequency of SNPs within the genome, there is a greater probability that a SNP will be linked to a genetic locus of interest than other types of genetic markers.

Disease associated SNPs can occur in coding and non-coding regions of the genome. When located in a coding region, the presence of the SNP can result in the production of a protein that is non-functional or has decreased function. More frequently, SNPs occur in non-coding regions. If the SNP occurs in a regulatory region, it may affect expression of the protein. For example, the presence of a SNP in a promoter region, may cause decreased expression of a protein. If the protein is involved in protecting the body against development of a pathological condition, this decreased expression can make the individual more susceptible to the condition.

Numerous methods exist for the detection of SNPs within a nucleotide sequence. A review of many of these methods can be found in Landegren et al., *Genome Res.*, 8:769-776, 1998. SNPs can be detected by restriction fragment length polymorphism (RFLP) (U.S. Patent Nos. 5,324,631; 5,645,995). RFLP analysis of the SNPs, however, is limited to cases where the SNP either creates or destroys a restriction enzyme cleavage site. SNPs can also be detected by direct sequencing of the nucleotide sequence of interest. Numerous assays based on hybridization have also been developed to detect SNPs. In addition, mismatch distinction by polymerases and ligases has also been used to detect SNPs.

There is growing recognition that SNPs can provide a powerful tool for the detection of individuals whose genetic make-up alters their susceptibility to certain diseases. There are four primary reasons why SNPs are especially suited for the identification of genotypes which predispose an individual to develop a disease condition. First, SNPs are by far the most prevalent type of polymorphism present in the genome and so are likely to be present in or near any locus of interest. Second, SNPs located in genes can be expected to directly affect protein structure or expression levels and so may serve not only as markers but as candidates for gene therapy treatments to cure or prevent a disease. Third, SNPs show greater genetic stability than repeated sequences and so are less likely to undergo changes which would complicate diagnosis. Fourth, the increasing efficiency of methods of detection of SNPs make them especially suitable for high throughput typing systems necessary to screen large populations.

SUMMARY

The present inventor has discovered novel single nucleotide polymorphisms (SNPs) associated with the development of various diseases, including hypertension (HTN), end stage renal disease due to hypertension (ESRD due to HTN), non-insulin dependent diabetes mellitus (NIDDM), end stage renal disease due to non-insulin dependent diabetes mellitus (ESRD due to NIDDM), lung cancer, breast cancer, prostate cancer, colon cancer, atherosclerotic peripheral vascular disease due to hypertension (ASPVD due to HTN), cerebrovascular accident due to hypertension (CVA due to HTN), cataracts due to hypertension (cataracts due to HTN), cardiomyopathy with hypertension (HTN CM), myocardial infarction due to hypertension (MI due to HTN), atherosclerotic peripheral vascular disease due to non-insulin dependent diabetes mellitus (ASPVD due to NIDDM), cerebrovascular accident due to non-insulin dependent diabetes mellitus (CVA

due to NIDDM), ischemic cardiomyopathy (Ischemic CM), ischemic cardiomyopathy with non-insulin dependent diabetes mellitus (Ischemic CM with NIDDM), myocardial infarction due to non-insulin dependent diabetes mellitus (MI due to NIDDM), atrial fibrillation without valvular disease (afib without valvular disease), alcohol abuse, anxiety,

5   asthma, chronic obstructive pulmonary disease (COPD), cholecystectomy, degenerative joint disease (DJD), end stage renal disease and frequent de-clots (ESRD and frequent de-clots), end stage renal disease due to focal segmental glomerular sclerosis (ESRD due to FSGS), end stage renal disease due to insulin dependent diabetes mellitus (ESRD due to IDDM), or seizure disorder. As such, this polymorphism provides a method for

10   diagnosing a genetic predisposition for the development of these diseases in individuals. Information obtained from the detection of SNPs associated with the development of these diseases is of great value in their treatment and prevention.

Accordingly, one aspect of the present invention provides a method for diagnosing a genetic predisposition for HTN, ESRD due to HTN, NIDDM, ESRD due to NIDDM,

15   lung cancer, breast cancer, prostate cancer, colon cancer, ASPVD due to HTN, CVA due to HTN, cataracts due to HTN, HTN CM, MI due to HTN, ASPVD due to NIDDM, CVA due to NIDDM, Ischemic CM, Ischemic CM with NIDDM, MI due to NIDDM, afib without valvular disease, alcohol abuse, anxiety, asthma, COPD, cholecystectomy, DJD, ESRD and frequent de-clots, ESRD due to FSGS, ESRD due to IDDM, or seizure disorder

20   in a subject, comprising obtaining a sample containing at least one polynucleotide from the subject, and analyzing the polynucleotide to detect a genetic polymorphism wherein the presence or absence of said genetic polymorphism is associated with an altered susceptibility to developing HTN, ESRD due to HTN, NIDDM, ESRD due to NIDDM, lung cancer, breast cancer, prostate cancer, colon cancer, ASPVD due to HTN, CVA due

25   to HTN, cataracts due to HTN, HTN CM, MI due to HTN, ASPVD due to NIDDM, CVA due to NIDDM, Ischemic CM, Ischemic CM with NIDDM, MI due to NIDDM, afib without valvular disease, alcohol abuse, anxiety, asthma, COPD, cholecystectomy, DJD, ESRD and frequent de-clots, ESRD due to FSGS, ESRD due to IDDM, or seizure disorder. In one embodiment, the polymorphism is located in the EDN-1 gene.

30   Another aspect of the present invention provides an isolated nucleic acid sequence comprising at least 10 contiguous nucleotides from SEQ ID NO: 1, or their complements, wherein the sequence contains at least one polymorphic site associated with a disease and in particular HTN, ESRD due to HTN, NIDDM, ESRD due to NIDDM, lung cancer, breast cancer, prostate cancer, colon cancer, ASPVD due to HTN, CVA due to HTN,

cataracts due to HTN, HTN CM, MI due to HTN, ASPVD due to NIDDM, CVA due to NIDDM, Ischemic CM, Ischemic CM with NIDDM, MI due to NIDDM, afib without valvular disease, alcohol abuse, anxiety, asthma, COPD, cholecystectomy, DJD, ESRD and frequent de-clots, ESRD due to FSGS, ESRD due to IDDM, or seizure disorder.

5   Yet another aspect of the invention is a kit for the detection of a polymorphism comprising, at a minimum, at least one polynucleotide of at least 10 contiguous nucleotides of SEQ ID NO: 1, or their complements, wherein the polynucleotide contains at least one polymorphic site associated with HTN, ESRD due to HTN, NIDDM, ESRD due to NIDDM, lung cancer, breast cancer, prostate cancer, colon cancer, ASPVD due to

10   HTN, CVA due to HTN, cataracts due to HTN, HTN CM, MI due to HTN, ASPVD due to NIDDM, CVA due to NIDDM, Ischemic CM, Ischemic CM with NIDDM, MI due to NIDDM, afib without valvular disease, alcohol abuse, anxiety, asthma, COPD, cholecystectomy, DJD, ESRD and frequent de-clots, ESRD due to FSGS, ESRD due to IDDM, or seizure disorder.

15   Yet another aspect of the invention provides a method for treating HTN, ESRD due to HTN, NIDDM, ESRD due to NIDDM, lung cancer, breast cancer, prostate cancer, colon cancer, ASPVD due to HTN, CVA due to HTN, cataracts due to HTN, HTN CM, MI due to HTN, ASPVD due to NIDDM, CVA due to NIDDM, Ischemic CM, Ischemic CM with NIDDM, MI due to NIDDM, afib without valvular disease, alcohol abuse,

20   anxiety, asthma, COPD, cholecystectomy, DJD, ESRD and frequent de-clots, ESRD due to FSGS, ESRD due to IDDM, or seizure disorder comprising, obtaining a sample of biological material containing at least one polynucleotide from the subject; analyzing the polynucleotide to detect the presence of at least one polymorphism associated with HTN, ESRD due to HTN, NIDDM, ESRD due to NIDDM, lung cancer, breast cancer, prostate

25   cancer, colon cancer, ASPVD due to HTN, CVA due to HTN, cataracts due to HTN, HTN CM, MI due to HTN, ASPVD due to NIDDM, CVA due to NIDDM, Ischemic CM, Ischemic CM with NIDDM, MI due to NIDDM, afib without valvular disease, alcohol abuse, anxiety, asthma, COPD, cholecystectomy, DJD, ESRD and frequent de-clots, ESRD due to FSGS, ESRD due to IDDM, or seizure disorder; and treating the subject in

30   such a way as to counteract the effect of any such polymorphism detected.

Still another aspect of the invention provides a method for the prophylactic treatment of a subject with a genetic predisposition to HTN, ESRD due to HTN, NIDDM, ESRD due to NIDDM, lung cancer, breast cancer, prostate cancer, colon cancer, ASPVD due to HTN, CVA due to HTN, cataracts due to HTN, HTN CM, MI due to HTN, ASPVD

due to NIDDM, CVA due to NIDDM, Ischemic CM, Ischemic CM with NIDDM, MI due to NIDDM, afib without valvular disease, alcohol abuse, anxiety, asthma, COPD, cholecystectomy, DJD, ESRD and frequent de-clots, ESRD due to FSGS, ESRD due to IDDM, or seizure disorder comprising, obtaining a sample of biological material

5   containing at least one polynucleotide from the subject; analyzing the polynucleotide to detect the presence of at least one polymorphism associated with HTN, ESRD due to HTN, NIDDM, ESRD due to NIDDM, lung cancer, breast cancer, prostate cancer, colon cancer, ASPVD due to HTN, CVA due to HTN, cataracts due to HTN, HTN CM, MI due to HTN, ASPVD due to NIDDM, CVA due to NIDDM, Ischemic CM, Ischemic CM with

10   NIDDM, MI due to NIDDM, afib without valvular disease, alcohol abuse, anxiety, asthma, COPD, cholecystectomy, DJD, ESRD and frequent de-clots, ESRD due to FSGS, ESRD due to IDDM, or seizure disorder; and treating the subject.

Further scope of the applicability of the present invention will become apparent from the detailed description and drawings provided below. It should be understood,

15   however, that the following detailed description and examples, while indicating preferred embodiments of the invention, are given by way of illustration only, since various changes and modifications within the spirit and scope of the invention will become apparent to those skilled in the art from the following detailed description.

20   **BRIEF DESCRIPTION OF THE DRAWINGS**

These and other features, aspects, and advantages of the present invention will become better understood with regard to the following description, appended claims, and accompanying drawings where:

Figure 1 shows SEQ ID NO: 1, the nucleotide sequence of the EDN-1 promoter

25   region as contained in GenBank Accession Number J05008.1. Position of the single nucleotide polymorphisms (SNPs) are here given according to the numbering scheme in GenBank Accession Number J05008.1. Thus, all nucleotides will be positively numbered, rather than bear negative numbers reflecting their position upstream from the RNA polymerase II binding site (a TATA box in about half of eukaryotic genes), the

30   transcription initiation site (a variable number of nucleotides downstream of, i.e. 3' to, the TATA box), the translation start site, or the first codon of the encoded protein (the "A" of the "ATG" codon for methionine, the first amino acid of every protein). Since not all genes are fully annotated, and not all promoter sequences contain elements far downstream such as the "ATG" encoding the first methionine in the translated protein, the

numbering system used in this patent application is less troublesome.

The various numbering systems can be easily interconverted, if desired. According to the annotation of Accession Number , the TATA box is located at position 3577. The first exon begins at position 3608. The position of the ATG codon for the first amino acid

5   (methionine) of the protein is at position 3876.

The first SNP mentioned below is located at position 2239 of Accession Number J05008.1. According to the annotation of Accession Number J05008.1, the transcription start site is position 3608. Therefore, the T2239->G SNP would be –1369 relative to the transcription start position. Further, according to the annotation of Accession Number

10   J05008.1, the position of the "A" of the ATG codon for the first amino acid (methionine) of the protein, i.e.- the translation start site, is at position 3876. The T2239→G SNP corresponds to –1637 with reference to the translation initiation site (the "A" of the first encoded "ATG").

The second SNP mentioned below (A2657→C) is located at position 2657

15   according to the numbering scheme of GenBank Accession Number J05008.1. Again, according to the annotation of Accession Number J05008.1, the transcription start site is position 3608. Therefore, the A2657->C SNP would be –951 relative to the transcription start position. Further, according to the annotation of Accession Number J05008.1, the position of the "A" of the ATG codon for the first amino acid (methionine) of the protein,

20   i.e.- the translation start site, is at position 3876. The A2657→C SNP corresponds to – 1219 with reference to the translation initiation site (the "A" of the first encoded "ATG").

DEFINITIONS

25   nt = nucleotide

bp = base pair

kb = kilobase; 1000 base pairs

ASPVD = atherosclerotic peripheral vascular disease

COPD = chronic obstructive pulmonary disease

30   CVA = cerebrovascular accident

DJD = degenerative joint disease, also know as osteoarthritis

DOL = dye-labeled oligonucleotide ligation assay

ESRD = end-stage renal disease

FSGS = focal segmental glomerular sclerosis

HTN = hypertension

MASDA = multiplexed allele-specific diagnostic assay

MADGE = microtiter array diagonal gel electrophoresis

MI = myocardial infarction

5   NIDDM = noninsulin-dependent diabetes mellitus

OLA = oligonucleotide ligation assay

PCR = polymerase chain reaction

RFLP = restriction fragment length polymorphism

SNP = single nucleotide polymorphism

10   "Polynucleotide" and "oligonucleotide" are used interchangeably and mean a linear polymer of at least 2 nucleotides joined together by phosphodiester bonds and may consist of either ribonucleotides or deoxyribonucleotides.

"Sequence" means the linear order in which monomers occur in a polymer, for example, the order of amino acids in a polypeptide or the order of nucleotides in a

15   polynucleotide.

"Polymorphism" refers to a set of genetic variants at a particular genetic locus among individuals in a population.

"Promoter" means a regulatory sequence of DNA that is involved in the binding of RNA polymerase to initiate transcription of a gene. A "gene" is a segment of DNA

20   involved in producing a peptide, polypeptide, or protein, including the coding region, non-coding regions preceding ("leader") and following ("trailer") coding region, as well as intervening non-coding sequences ("introns") between individual coding segments ("exons"). A promoter is herein considered as a part of the corresponding gene. Coding refers to the representation of amino acids, start and stop signals in a three base "triplet"

25   code. Promoters are often upstream ("5' to") the transcription initiation site of the gene.

"Gene therapy" means the introduction of a functional gene or genes from some source by any suitable method into a living cell to correct for a genetic defect.

"Reference allele" or "reference type" means the allele designated in the Gen Bank sequence listing for a given gene, in this case Gen Bank Accession Number J05008.1 for

30   the endothelin-1 gene.

"Genetic variant" or "variant" means a specific genetic variant which is present at a particular genetic locus in at least one individual in a population and that differs from the reference type.

As used herein the terms "patient" and "subject" are not limited to human beings, but are intended to include all vertebrate animals in addition to human beings.

As used herein the terms "genetic predisposition", "genetic susceptibility" and "susceptibility" all refer to the likelihood that an individual subject will develop a
5 particular disease, condition or disorder. For example, a subject with an increased susceptibility or predisposition will be more likely than average to develop a disease, while a subject with a decreased predisposition will be less likely than average to develop the disease. A genetic variant is associated with an altered susceptibility or predisposition if the allele frequency of the genetic variant in a population or subpopulation with a
10 disease, condition or disorder varies from its allele frequency in the population without the disease, condition or disorder (control population) or a control sequence (reference type) by at least 1%, preferably by at least 2%, more preferably by at least 4% and more preferably still by at least 8%. Alternatively, an odds ratio of 1.5 was chosen as the threshold of significance based on the recommendation of Austin et al. in *Epidemiol. Rev.*,
15 16:65-76, 1994. "[E]pidemiology in general and case-control studies in particular are not well suited for detecting weak associations (odds ratios < 1.5)." *Id.* at 66.

As used herein "isolated nucleic acid" means a species of the invention that is the predominate species present (i.e., on a molar basis it is more abundant than any other individual species in the composition). Preferably, an isolated nucleic acid comprises at
20 least about 50, 80 or 90 percent (on a molar basis) of all macromolecular species present. Most preferably, the object species is purified to essential homogeneity (contaminant species cannot be detected in the composition by conventional detection methods).

As used herein, "allele frequency" means the frequency that a given allele appears in a population.

25

DETAILED DESCRIPTION

All publications, patents, patent applications and other references cited in this application are herein incorporated by reference in their entirety as if each individual publication, patent, patent application or other reference were specifically and individually
30 indicated to be incorporated by reference.

Novel Polymorphisms

The present application provides single nucleotide polymorphisms (SNPs) in a gene associated with HTN, ESRD due to HTN, NIDDM, ESRD due to NIDDM, lung

cancer, breast cancer, prostate cancer, colon cancer, ASPVD due to HTN, CVA due to HTN, cataracts due to HTN, HTN CM, MI due to HTN, ASPVD due to NIDDM, CVA due to NIDDM, Ischemic CM, Ischemic CM with NIDDM, MI due to NIDDM, afib without valvular disease, alcohol abuse, anxiety, asthma, COPD, cholecystectomy, DJD, ESRD and frequent de-clots, ESRD due to FSGS, ESRD due to IDDM, or seizure disorder. The first polymorphism is a T to G transversion at position 2239 and the second polymorphism is an A to C substitution at position 2657, both of the EDN-1 promoter.

Preparation of Samples

The presence of genetic variants in the above genes or their control regions, or in any other genes that may affect susceptibility to disease is determined by screening nucleic acid sequences from a population of individuals for such variants. The population is preferably comprised of some individuals with the disease of interest, so that any genetic variants that are found can be correlated with disease. The population is also preferably comprised of some individuals that have known risk for the disease. The population should preferably be large enough to have a reasonable chance of finding individuals with the sought-after genetic variant. As the size of the population increases, the ability to find significant correlations between a particular genetic variant and susceptibility to disease also increases.

The nucleic acid sequence can be DNA or RNA. For the assay of genomic DNA, virtually any biological sample containing genomic DNA (e.g., not pure red blood cells) can be used. For example, and without limitation, genomic DNA can be conveniently obtained from whole blood, semen, saliva, tears, urine, fecal material, sweat, buccal cells, skin or hair. For assays using cDNA or mRNA, the target nucleic acid must be obtained from cells or tissues that express the target sequence. One preferred source and quantity of DNA is 10 to 30 ml of anticoagulated whole blood, since enough DNA can be extracted from leukocytes in such a sample to perform many repetitions of the analysis contemplated herein.

Many of the methods described herein require the amplification of DNA from target samples. This can be accomplished by any method known in the art but preferably is by the polymerase chain reaction (PCR). Optimization of conditions for conducting PCR must be determined for each reaction and can be accomplished without undue experimentation by one of ordinary skill in the art. In general, methods for conducting PCR can be found in U.S. Patent Nos 4,965,188, 4,800,159, 4,683,202, and 4,683,195;

Ausbel et al., eds., *Short Protocols in Molecular Biology*, 3rd ed., Wiley, 1995; and Innis et al., eds., *PCR Protocols*, Academic Press, 1990.

Other amplification methods include the ligase chain reaction (LCR) (see, Wu and Wallace, *Genomics*, 4:560-569, 1989; Landegren et al., *Science*, 241:1077-1080, 1988), transcription amplification (Kwoh et al., *Proc. Natl. Acad. Sci. USA*, 86:1173-1177, 1989), self-sustained sequence replication (Guatelli et al., *Proc. Natl. Acad. Sci. USA*, 87:1874-1878, 1990), and nucleic acid based sequence amplification (NASBA). The latter two amplification methods involve isothermal reactions based on isothermal transcription, which produces both single stranded RNA (ssRNA) and double stranded DNA (dsDNA) as the amplification products in a ratio of about 30 or 100 to 1, respectively.

Detection of Polymorphisms

Detection of Unknown Polymorphisms

Two types of detection are contemplated within the present invention. The first type involves detection of unknown SNPs by comparing nucleotide target sequences from individuals in order to detect sites of polymorphism. If the most common sequence of the target nucleotide sequence is not known, it can be determined by analyzing individual humans, animals or plants with the greatest diversity possible. Additionally the frequency of sequences found in subpopulations characterized by such factors as geography or gender can be determined.

The presence of genetic variants and in particular SNPs is determined by screening the DNA and/or RNA of a population of individuals for such variants. If it is desired to detect variants associated with a particular disease or pathology, the population is preferably comprised of some individuals with the disease or pathology, so that any genetic variants that are found can be correlated with the disease of interest. It is also preferable that the population be composed of individuals with known risk factors for the disease. The populations should preferably be large enough to have a reasonable chance to find correlations between a particular genetic variant and susceptibility to the disease of interest. In addition, the allele frequency of the genetic variant in a population or subpopulation with the disease or pathology should vary from its allele frequency in the population without the disease or pathology (control population) or the control sequence (reference type) by at least 1%, preferably by at least 2%, more preferably by at least 4% and more preferably still by at least 8%.

Determination of unknown genetic variants, and in particular SNPs, within a particular nucleotide sequence among a population may be determined by any method known in the art, for example and without limitation, direct sequencing, restriction length fragment polymorphism (RFLP), single-strand conformational analysis (SSCA), denaturing gradient gel electrophoresis (DGGE), heteroduplex analysis (HET), chemical cleavage analysis (CCM) and ribonuclease cleavage.

Methods for direct sequencing of nucleotide sequences are well known to those skilled in the art and can be found for example in Ausubel et al., eds., *Short Protocols in Molecular Biology*, 3rd ed., Wiley, 1995 and Sambrook et al., *Molecular Cloning*, 2nd ed., Chap. 13, Cold Spring Harbor Laboratory Press, 1989. Sequencing can be carried out by any suitable method, for example, dideoxy sequencing (Sanger et al., *Proc. Natl. Acad. Sci. USA*, 74:5463-5467, 1977), chemical sequencing (Maxam and Gilbert, *Proc. Natl. Acad. Sci. USA*, 74:560-564, 1977) or variations thereof. Direct sequencing has the advantage of determining variation in any base pair of a particular sequence.

RFLP analysis (see, e.g. U.S. Patents No. 5,324,631 and 5,645,995) is useful for detecting the presence of genetic variants at a locus in a population when the variants differ in the size of a probed restriction fragment within the locus, such that the difference between the variants can be visualized by electrophoresis. Such differences will occur when a variant creates or eliminates a restriction site within the probed fragment. RFLP analysis is also useful for detecting a large insertion or deletion within the probed fragment. Thus, RFLP analysis is useful for detecting, e.g., an *Alu* sequence insertion or deletion in a probed DNA segment.

Single-strand conformational polymorphisms (SSCPs) can be detected in <220 bp PCR amplicons with high sensitivity (Orita et al, *Proc. Natl. Acad. Sci. USA*, 86:2766-2770, 1989; Warren et al., In: *Current Protocols in Human Genetics*, Dracopoli et al., eds, Wiley, 1994, 7.4.1-7.4.6.). Double strands are first heat-denatured. The single strands are then subjected to polyacrylamide gel electrophoresis under non-denaturing conditions at constant temperature (i.e., low voltage and long run times) at two different temperatures, typically 4-10°C and 23°C (room temperature). At low temperatures (4-10°C), the secondary structure of short single strands (degree of intrachain hairpin formation) is sensitive to even single nucleotide changes, and can be detected as a large change in electrophoretic mobility. The method is empirical, but highly reproducible, suggesting the existence of a very limited number of folding pathways for short DNA strands at the

critical temperature. Polymorphisms appear as new banding patterns when the gel is stained.

Denaturing gradient gel electrophoresis (DGGE) can detect single base mutations based on differences in migration between homo- and heteroduplexes (Myers et al., *Nature*, 313:495-498, 1985). The DNA sample to be tested is hybridized to a labeled reference type probe. The duplexes formed are then subjected to electrophoresis through a polyacrylamide gel that contains a gradient of DNA denaturant parallel to the direction of electrophoresis. Heteroduplexes formed due to single base variations are detected on the basis of differences in migration between the heteroduplexes and the homoduplexes formed.

In heteroduplex analysis (HET) (Keen et al., *Trends Genet.*7:5, 1991), genomic DNA is amplified by the polymerase chain reaction followed by an additional denaturing step which increases the chance of heteroduplex formation in heterozygous individuals. The PCR products are then separated on Hydrolink gels where the presence of the heteroduplex is observed as an additional band.

Chemical cleavage analysis (CCM) is based on the chemical reactivity of thymine (T) when mismatched with cytosine, guanine or thymine and the chemical reactivity of cytosine (C) when mismatched with thymine, adenine or cytosine (Cotton et al., *Proc. Natl. Acad. Sci. USA*, 85:4397-4401, 1988). Duplex DNA formed by hybridization of a reference type probe with the DNA to be examined, is treated with osmium tetroxide for T and C mismatches and hydroxylamine for C mismatches. T and C mismatched bases that have reacted with the hydroxylamine or osmium tetroxide are then cleaved with piperidine. The cleavage products are then analyzed by gel electrophoresis.

Ribonuclease cleavage involves enzymatic cleavage of RNA at a single base mismatch in an RNA:DNA hybrid (Myers et al., *Science* 230:1242-1246, 1985). A ^{32}P labeled RNA probe complementary to the reference type DNA is annealed to the test DNA and then treated with ribonuclease A. If a mismatch occurs, ribonuclease A will cleave the RNA probe and the location of the mismatch can then be determined by size analysis of the cleavage products following gel electrophoresis.

Detection of Known Polymorphisms

The second type of polymorphism detection involves determining which form of a known polymorphism is present in individuals for diagnostic or epidemiological purposes.

In addition to the already discussed methods for detection of polymorphisms, several methods have been developed to detect known SNPs. Many of these assays have been reviewed by Landegren et al., *Genome Res.*, 8:769-776, 1998, and will only be briefly reviewed here.

One type of assay has been termed an array hybridization assay, an example of which is the multiplexed allele-specific diagnostic assay (MASDA) (U.S. Patent No. 5,834,181; Shuber et al., *Hum. Molec. Genet.*, 6:337-347, 1997). In MASDA, samples from multiplex PCR are immobilized on a solid support. A single hybridization is conducted with a pool of labeled allele specific oligonucleotides (ASO). Any ASOs that hybridize to the samples are removed from the pool of ASOs. The support is then washed to remove unhybridized ASOs remaining in the pool. Labeled ASOs remaining on the support are detected and eluted from the support. The eluted ASOs are then sequenced to determine the mutation present.

Two assays depend on hybridization-based allele-discrimination during PCR. The TaqMan assay (U.S. Patent No. 5,962,233; Livak et al., *Nature Genet.*, 9:341-342, 1995) uses allele specific (ASO) probes with a donor dye on one end and an acceptor dye on the other end, such that the dye pair interact via fluorescence resonance energy transfer (FRET). A target sequence is amplified by PCR modified to include the addition of the labeled ASO probe. The PCR conditions are adjusted so that a single nucleotide difference will effect binding of the probe. Due to the 5′ nuclease activity of the *Taq* polymerase enzyme, a perfectly complementary probe is cleaved during the PCR while a probe with a single mismatched base is not cleaved. Cleavage of the probe dissociates the donor dye from the quenching acceptor dye, greatly increasing the donor fluorescence.

An alternative to the TaqMan assay is the molecular beacons assay (U.S. Patent No. 5,925,517; Tyagi et al., *Nature Biotech.*, 16:49-53, 1998). In the molecular beacons

method for real time detection of the presence of target sequences or can be used after amplification.

High throughput screening for SNPs that affect restriction sites can be achieved by Microtiter Array Diagonal Gel Electrophoresis (MADGE) (Day and Humphries, *Anal. Biochem.*, 222:389-395, 1994). In this assay restriction fragment digested PCR products are loaded onto stackable horizontal gels with the wells arrayed in a microtiter format. During electrophoresis, the electric field is applied at an angle relative to the columns and rows of the wells allowing products from a large number of reactions to be resolved.

Additional assays for SNPs depend on mismatch distinction by polymerases and ligases. The polymerization step in PCR places high stringency requirements on correct base pairing of the 3' end of the hybridizing primers. This has allowed the use of PCR for the rapid detection of single base changes in DNA by using specifically designed oligonucleotides in a method variously called PCR amplification of specific alleles (PASA) (Sommer et al., *Mayo Clin. Proc.*, 64:1361-1372, 1989; Sarker et al., *Anal. Biochem.*, 1990), allele-specific amplification (ASA), allele-specific PCR, and amplification refractory mutation system (ARMS) (Newton et al., *Nuc. Acids Res.*, 1989; Nichols et al., *Genomics*, 1989; Wu et al., *Proc. Natl. Acad. Sci. USA*, 1989). In these methods, an oligonucleotide primer is designed that perfectly matches one allele but mismatches the other allele at or near the 3' end. This results in the preferential amplification of one allele over the other. By using three primers that produce two differently sized products, it can be determined whether an individual is homozygous or heterozygous for the mutation (Dutton and Sommer, *BioTechniques*, 11:700-702, 1991). In another method, termed bi-PASA, four primers are used; two outer primers that bind at different distances from the site of the SNP and two allele specific inner primers (Liu et al., *Genome Res.*, 7:389-398, 1997). Each of the inner primers has a non-complementary 5' end and form a mismatch near the 3' end if the proper allele is not present. Using this system, zygosity is determined based on the size and number of PCR products produced.

The joining by DNA ligases of two oligonucleotides hybridized to a target DNA sequence is quite sensitive to mismatches close to the ligation site, especially at the 3' end. This sensitivity has been utilized in the oligonucleotide ligation assay (Landegren et al., *Science*, 241:1077-1080, 1988) and the ligase chain reaction (LCR; Barany, *Proc. Natl. Acad. Sci. USA*, 88:189-193, 1991). In OLA, the sequence surrounding the SNP is first amplified by PCR, whereas in LCR, genomic DNA can be used as a template.

In one method for mass screening for SNPs based on the OLA, amplified DNA templates are analyzed for their ability to serve as templates for ligation reactions between labeled oligonucleotide probes (Samotiaki et al., *Genomics*, 20:238-242, 1994). In this assay, two allele-specific probes labeled with either of two lanthanide labels (europium or terbium) compete for ligation to a third biotin labeled phosphorylated oligonucleotide and the signals from the allele specific oligonucleotides are compared by time-resolved fluorescence. After ligation, the oligonucleotides are collected on an avidin-coated 96-pin capture manifold. The collected oligonucleotides are then transferred to microtiter wells in which the europium and terbium ions are released. The fluorescence from the europium ions is determined for each well, followed by measurement of the terbium fluorescence.

In alternative gel-based OLA assays, numerous SNPs can be detected simultaneously using multiplex PCR and multiplex ligation (U.S. Patent No. 5,830,711; Day et al., *Genomics*, 29:152-162, 1995; Grossman et al., *Nuc. Acids Res.*, 22:4527-4534, 1994). In these assays, allele specific oligonucleotides with different markers, for example, fluorescent dyes, are used. The ligation products are then analyzed together by electrophoresis on an automatic DNA sequencer distinguishing markers by size and alleles by fluorescence. In the assay by Grossman et al., 1994, mobility is further modified by the presence of a non-nucleotide mobility modifier on one of the oligonucleotides.

A further modification of the ligation assay has been termed the dye-labeled oligonucleotide ligation (DOL) assay (U.S. Patent No. 5,945,283; Chen et al., *Genome Res.*, 8:549-556, 1998). DOL combines PCR and the oligonucleotide ligation reaction in a two-stage thermal cycling sequence with fluorescence resonance energy transfer (FRET) detection. In the assay, labeled ligation oligonucleotides are designed to have annealing temperatures lower than those of the amplification primers. After amplification, the temperature is lowered to a temperature where the ligation oligonucleotides can anneal and be ligated together. This assay requires the use of a thermostable ligase and a thermostable DNA polymerase without 5' nuclease activity. Because FRET occurs only when the donor and acceptor dyes are in close proximity, ligation is inferred by the change in fluorescence.

In another method for the detection of SNPs termed minisequencing, the target-dependent addition by a polymerase of a specific nucleotide immediately downstream (3') to a single primer is used to determine which allele is present (U.S Patent No. 5,846,710). Using this method, several SNPs can be analyzed in parallel by separating locus specific

primers on the basis of size via electrophoresis and determining allele specific incorporation using labeled nucleotides.

Determination of individual SNPs using solid phase minisequencing has been described by Syvanen et al., *Am. J. Hum. Genet.*, 52:46-59, 1993. In this method the sequence including the polymorphic site is amplified by PCR using one amplification primer which is biotinylated on its 5' end. The biotinylated PCR products are captured in streptavidin-coated microtitration wells, the wells washed, and the captured PCR products denatured. A sequencing primer is then added whose 3' end binds immediately prior to the polymorphic site, and the primer is elongated by a DNA polymerase with one single labeled dNTP complementary to the nucleotide at the polymorphic site. After the elongation reaction, the sequencing primer is released and the presence of the labeled nucleotide detected. Alternatively, dye labeled dideoxynucleoside triphosphates (ddNTPs) can be used in the elongation reaction (U.S. Patent No. 5,888,819; Shumaker et al., *Human Mut.*, 7:346-354, 1996). In this method, incorporation of the ddNTP is determined using an automatic gel sequencer.

Minisequencing has also been adapted for use with microarrays (Shumaker et al., *Human Mut.*, 7:346-354, 1996). In this case, elongation (extension) primers are attached to a solid support such as a glass slide. Methods for construction of oligonucleotide arrays are well known to those of ordinary skill in the art and can be found, for example, in *Nature Genetics*, Suppl., Vol. 21, January, 1999. PCR products are spotted on the array and allowed to anneal. The extension (elongation) reaction is carried out using a polymerase, a labeled dNTP and noncompeting ddNTPs. Incorporation of the labeled dNTP is then detected by the appropriate means. In a variation of this method suitable for use with multiplex PCR, extension is accomplished with the use of the appropriate labeled ddNTP and unlabeled ddNTPs (Pastinen et al., *Genome Res.*, 7:606-614, 1997).

Solid phase minisequencing has also been used to detect multiple polymorphic nucleotides from different templates in an undivided sample (Pastinen et al., *Clin. Chem.*, 42:1391-1397, 1996). In this method, biotinylated PCR products are captured on the avidin-coated manifold support and rendered single stranded by alkaline treatment. The manifold is then placed serially in four reaction mixtures containing extension primers of varying lengths, a DNA polymerase and a labeled ddNTP, and the extension reaction allowed to proceed. The manifolds are inserted into the slots of a gel containing formamide which releases the extended primers from the template. The extended primers are then identified by size and fluorescence on a sequencing instrument.

Fluorescence resonance energy transfer (FRET) has been used in combination with minisequencing to detect SNPs (U.S. Patent No. 5,945,283; Chen et al., *Proc. Natl. Acad. Sci. USA*, 94:10756-10761, 1997). In this method, the extension primers are labeled with a fluorescent dye, for example fluorescein. The ddNTPs used in primer extension are labeled with an appropriate FRET dye. Incorporation of the ddNTPs is determined by changes in fluorescence intensities.

The above discussion of methods for the detection of SNPs is exemplary only and is not intended to be exhaustive. Those of ordinary skill in the art will be able to envision other methods for detection of SNPs that are within the scope and spirit of the present invention.

In one embodiment the present invention provides a method for diagnosing a genetic predisposition for a disease. In this method, a biological sample is obtained from a subject. The subject can be a human being or any vertebrate animal. The biological sample must contain polynucleotides and preferably genomic DNA. Samples that do not contain genomic DNA, for example, pure samples of mammalian red blood cells, are not suitable for use in the method. The form of the polynucleotide is not critically important such that the use of DNA, cDNA, RNA or mRNA is contemplated within the scope of the method. The polynucleotide is then analyzed to detect the presence of a genetic variant where such variant is associated with an increased risk of developing a disease, condition or disorder, and in particular HTN, ESRD due to HTN, NIDDM, ESRD due to NIDDM, lung cancer, breast cancer, prostate cancer, colon cancer, ASPVD due to HTN, CVA due to HTN, cataracts due to HTN, HTN CM, MI due to HTN, ASPVD due to NIDDM, CVA due to NIDDM, Ischemic CM, Ischemic CM with NIDDM, MI due to NIDDM, afib without valvular disease, alcohol abuse, anxiety, asthma, COPD, cholecystectomy, DJD, ESRD and frequent de-clots, ESRD due to FSGS, ESRD due to IDDM, or seizure disorder. In one embodiment, the genetic variant is at one of the polymorphic sites contained in Table 17. In another embodiment, the genetic variant is one of the variants contained in Table 17 or the complement of any of the variants contained in Table 17. Any method capable of detecting a genetic variant, including any of the methods previously discussed, can be used. Suitable methods include, but are not limited to, those methods based on sequencing, mini sequencing, hybridization, restriction fragment analysis, oligonucleotide ligation, or allele specific PCR.

The present invention is also directed to an isolated nucleic acid sequence of at least 10 contiguous nucleotides from SEQ ID NO: 1, or the complements of SEQ ID NO:

1. In one preferred embodiment, the sequence contains at least one polymorphic site associated with a disease, and in particular HTN, ESRD due to HTN, NIDDM, ESRD due to NIDDM, lung cancer, breast cancer, prostate cancer, colon cancer, ASPVD due to HTN, CVA due to HTN, cataracts due to HTN, HTN CM, MI due to HTN, ASPVD due to NIDDM, CVA due to NIDDM, Ischemic CM, Ischemic CM with NIDDM, MI due to NIDDM, afib without valvular disease, alcohol abuse, anxiety, asthma, COPD, cholecystectomy, DJD, ESRD and frequent de-clots, ESRD due to FSGS, ESRD due to IDDM, or seizure disorder. In one embodiment, the genetic variant is at one of the polymorphic sites contained in Table 17. In another embodiment, the genetic variant is one of the variants contained in Table 17 or the complement of any of the variants contained in Table 17. In yet another embodiment, the polymorphic site, which may or may not also include a genetic variant, is located at the 3' end of the polynucleotide. In still another embodiment, the polynucleotide further contains a detectable marker. Suitable markers include, but are not limited to, radioactive labels, such as radionuclides, fluorophores or fluorochromes, peptides, enzymes, antigens, antibodies, vitamins or steroids.

The present invention also includes kits for the detection of polymorphisms associated with diseases, conditions or disorders, and in particular HTN, ESRD due to HTN, NIDDM, ESRD due to NIDDM, lung cancer, breast cancer, prostate cancer, colon cancer, ASPVD due to HTN, CVA due to HTN, cataracts due to HTN, HTN CM, MI due to HTN, ASPVD due to NIDDM, CVA due to NIDDM, Ischemic CM, Ischemic CM with NIDDM, MI due to NIDDM, afib without valvular disease, alcohol abuse, anxiety, asthma, COPD, cholecystectomy, DJD, ESRD and frequent de-clots, ESRD due to FSGS, ESRD due to IDDM, or seizure disorder. The kits contain, at a minimum, at least one polynucleotide of at least 10 contiguous nucleotides of SEQ ID NO 1, or the complements of SEQ ID NO: 1. In one embodiment, the genetic variant is at one of the polymorphic sites contained in Table 17. Alternatively the 3' end of the polynucleotide is immediately 5' to a polymorphic site, preferably a polymorphic site selected from the sites in Table 17. In another embodiment, the genetic variant is one of the variants contained in Table 17 or the complement of any of the variants contained in Table 17. In still another embodiment, the genetic variant is located at the 3' end of the polynucleotide. In yet another embodiment, the polynucleotide of the kit contains a detectable label. Suitable labels include, but are not limited to, radioactive labels, such as radionuclides, fluorophores or fluorochromes, peptides, enzymes, antigens, antibodies, vitamins or steroids.

In addition, the kit may also contain additional materials for detection of the polymorphisms. For example, and without limitation, the kits may contain buffer solutions, enzymes, nucleotide triphosphates, and other reagents and materials necessary for the detection of genetic polymorphisms. Additionally, the kits may contain

5   instructions for conducting analyses of samples for the presence of polymorphisms and for interpreting the results obtained.

In yet another embodiment the present invention provides a method for designing a treatment regime for a patient having a disease, condition or disorder and in particular HTN, ESRD due to HTN, NIDDM, ESRD due to NIDDM, lung cancer, breast cancer,

10   prostate cancer, colon cancer, ASPVD due to HTN, CVA due to HTN, cataracts due to HTN, HTN CM, MI due to HTN, ASPVD due to NIDDM, CVA due to NIDDM, Ischemic CM, Ischemic CM with NIDDM, MI due to NIDDM, afib without valvular disease, alcohol abuse, anxiety, asthma, COPD, cholecystectomy, DJD, ESRD and frequent de-clots, ESRD due to FSGS, ESRD due to IDDM, or seizure disorder caused either directly

15   or indirectly by the presence of one or more single nucleotide polymorphisms. In this method genetic material from a patient, for example, DNA, cDNA, RNA or mRNA is screened for the presence of one or more SNPs associated with the disease of interest. Depending on the type and location of the SNP, a treatment regime is designed to counteract the effect of the SNP.

20   Alternatively, information gained from analyzing genetic material for the presence of polymorphisms can be used to design treatment regimes involving gene therapy. For example, detection of a polymorphism that either affects the expression of a gene or results in the production of a mutant protein can be used to design an artificial gene to aid in the production of normal, wild type protein or help restore normal gene expression.

25   Methods for the construction of polynucleotide sequences encoding proteins and their associated regulatory elements are well know to those of ordinary skill in the art. Once designed, the gene can be placed in the individual by any suitable means known in the art (*Gene Therapy Technologies, Applications and Regulations*, Meager, ed., Wiley, 1999; *Gene Therapy: Principles and Applications*, Blankenstein, ed., Birkhauser Verlag, 1999;

30   Jain, *Textbook of Gene Therapy*, Hogrefe and Huber, 1998).

The present invention is also useful in designing prophylactic treatment regimes for patients determined to have an increased susceptibility to a disease, condition or disorder, and in particular HTN, ESRD due to HTN, NIDDM, ESRD due to NIDDM, lung cancer, breast cancer, prostate cancer, colon cancer, ASPVD due to HTN, CVA due to

HTN, cataracts due to HTN, HTN CM, MI due to HTN, ASPVD due to NIDDM, CVA due to NIDDM, Ischemic CM, Ischemic CM with NIDDM, MI due to NIDDM, afib without valvular disease, alcohol abuse, anxiety, asthma, COPD, cholecystectomy, DJD, ESRD and frequent de-clots, ESRD due to FSGS, ESRD due to IDDM, or seizure disorder

5   due to the presence of one or more single nucleotide polymorphisms. In this embodiment, genetic material, such as DNA, cDNA, RNA or mRNA, is obtained from a patient and screened for the presence of one or more SNPs associated either directly or indirectly to a disease, condition, disorder or other pathological condition. Based on this information, a treatment regime can be designed to decrease the risk of the patient developing the

10   disease. Such treatment can include, but is not limited to, surgery, the administration of pharmaceutical compounds or nutritional supplements, and behavioral changes such as improved diet, increased exercise, reduced alcohol intake, smoking cessation, etc.

EXAMPLES

15   The positions of the single nucleotide polymorphisms (SNPs) are given according to the numbering scheme in GenBank Accession Number J05008.1. Thus, all nucleotides will be positively numbered, rather than bear negative numbers reflecting their position upstream from the transcription initiation site, a scheme often used for promoters. The two numbering systems can be easily interconverted, if necessary. GenBank sequences

20   can be found at http://www.ncbi.nlm.nih.gov/

In the following examples, SNPs are written as "reference sequence nucleotide" → "variant nucleotide." Changes in nucleotide sequences are indicated in bold print. The standard nucleotide abbreviations are used in which A=adenine, C=cytosine, G=guanine, T=thymine, M=A or C, R=A or G, W=A or T, S=C or G, Y=C or T, K=G or T, V=A or C

25   or G, H=A or C or T; D=A or G or T; B=C or G or T; N= A or C or G or T.

Example 1
Detection of Novel Polymorphisms by Direct Sequencing of
Leukocyte Genomic DNA

30   Leukocytes were obtained from human whole blood collected with EDTA as an anticoagulant. Blood was obtained from a group of black men, black women, white men, and white women without any known disease. Blood was also obtained from individuals with HTN, ESRD due to HTN, NIDDM, ESRD due to NIDDM, lung cancer, breast cancer, prostate cancer, colon cancer, ASPVD due to HTN, CVA due to HTN, cataracts

due to HTN, HTN CM, MI due to HTN, ASPVD due to NIDDM, CVA due to NIDDM,
Ischemic CM, Ischemic CM with NIDDM, MI due to NIDDM, afib without valvular
disease, alcohol abuse, anxiety, asthma, COPD, cholecystectomy, DJD, ESRD and
frequent de-clots, ESRD due to FSGS, ESRD due to IDDM, or seizure disorder as
5   indicated in the tables below.

Genomic DNA was purified from the collected leukocytes using standard protocols
well known to those of ordinary skill in the art of molecular biology (Ausubel et al., *Short
Protocol in Molecular Biology*, 3rd ed., John Wiley and Sons, 1995; Sambrook et al.,
Molecular Cloning, Cold Spring Harbor Laboratory Press, 1989; and Davis et al., *Basic
10   Methods in Molecular Biology*, Elsevier Science Publishing, 1986). One hundred
nanograms of purified genomic DNA were used in each PCR reaction.

Standard PCR reaction conditions were used. Methods for conducting PCR are
well known in the art and can be found, for example, in U.S. Patent Nos 4,965,188,
4,800,159, 4,683,202, and 4,683,195; Ausbel et al., eds., *Short Protocols in Molecular
15   Biology*, 3rd ed., Wiley, 1995; and Innis et al., eds., *PCR Protocols*, Academic Press, 1990.

The first SNP T2239→G can be identified by PCR amplification of a specific
region of the endothelin-1 promoter. The sequence of the sense primer was 5'-CTC CAT
CCC CAG AAA AAC TG-3', corresponding to nucleotides 2113 to 2132, inclusive.
(SEQ ID NO: 2). The sequence of the anti-sense primer is 5'-AAG GAA GGT GGT GCT
20   GAG AA-3' corresponding to nucleotides 2490 to 2509, inclusive. (SEQ ID NO: 3). The
PCR product spanned positions 2113 to 2509, inclusive, of the EDN1 gene.

The second SNP A2657→C can be identified by PCR amplification of a specific
region of the endothelin-1 promoter. The sense primer was 5'- GGG GGA TTT CAA
GGT TAG AT -3' (SEQ ID NO: 4). The anti-sense primer was 5'- GAG AAG CCC CGA
25   TAA GTT CTT T -3' (SEQ ID NO: 5). The PCR product thus produced spanned
positions 2390 to 2924 of the human EDN-1 gene (SEQ ID NO: 1).

The PCR reaction contained a total volume of 20 microliters (μl), consisting of 10
μl of a premade PCR reaction mix (Sigma "JumpStart Ready Mix with RED Taq
Polymerase"). Primers at 10 μM were diluted to a final concentration of 0.3 μM in the
30   PCR reaction mix. Approximately 25 ng of template leukocyte genomic DNA was used
for each PCR amplification. After an initial 5 minutes denaturation at 94°C, 35 cycles
were performed consisting of 45 seconds of denaturation at 94°C, 45 seconds of
hybridization at 62°C, 45 seconds of extension at 72°C, followed by a final extension step
of 10 minutes at 72°C.

Post-PCR clean-up was performed as follows. PCR reactions were cleaned to remove unwanted primer and other impurities such as salts, enzymes, and unincorporated nucleotides that could inhibit sequencing. One of the following clean-up kits was used: Qiaquick-96 PCR Purification Kit (Qiagen) or Multiscreen-PCR Plates (Millipore,

5   discussed below).

When using the Qiaquick protocol, PCR samples were added to the 96-well Qiaquick silica-gel membrane plate and a chaotropic salt, supplied as "PB Buffer," was then added to each well. The PB Buffer caused the DNA to bind to the membrane. The plate was put onto the Qiagen vacuum manifold and vacuum was applied to the plate in

10   order to pull sample and PB Buffer through the membrane. The filtrate was discarded. Next, the samples were washed twice using "PE Buffer." Vacuum pressure was applied between each step to remove the buffer. Filtrate was similarly discarded after each wash. After the last PE Buffer wash, maximum vacuum pressure was applied to the membrane plate to generate maximum airflow through the membrane in order to evaporate residual

15   ethanol left from the PE Buffer. The clean PCR product was then eluted from the filter using "EB Buffer." The filtrate contained the cleaned PCR product and was collected. All buffers were supplied as part of the Qiaquick-96 PCR Purification Kit. The vacuum manifold was also purchased from Qiagen for exclusive use with the Qiaquick-96 Purification Kit.

20   When using the Millipore Multiscreen-PCR Plates, PCR samples were loaded into the wells of the Multiscreen-PCR Plate and the plate was then placed on a Millipore vacuum manifold. Vacuum pressure was applied for 10 minutes, and the filtrate was discarded. The plate was then removed from the vacuum manifold and 100 µl of Milli-Q water was added to each well to rehydrate the DNA samples. After shaking on a plate

25   shaker for 5 minutes, the plate was replaced on the manifold and vacuum pressure was applied for 5 minutes. The filtrate was again discarded. The plate was removed and 60 µl Milli-Q water was added to each well to again rehydrate the DNA samples. After shaking on a plate shaker for 10 minutes, the 60 µl of cleaned PCR product was transferred from the Multiscreen-PCR plate to another 96-well plate by pipetting. The Millipore vacuum

30   manifold was purchased from Millipore for exclusive use with the Multiscreen-PCR plates.

Cycle sequencing was performed on the clean PCR product using an ABI Prism Big Dye Terminator Cycle Sequencing Ready Reaction kit (Perkin-Elmer). For a total

volume of 20 µl, the following reagents were added to each well of a 96-well plate: 2.0 µl Terminator Ready Reaction mix, 3.0 µl 5X Sequencing Buffer (ABI), 5-10 µl template (30-90 ng double stranded DNA), 3.2 pM primer (primer used was the forward primer from the PCR reaction), and Milli-Q water to 20 µl total volume. The reaction plate was placed into a Hybaid thermal cycler block and programmed as follows: X 1 cycle: 1 degree/sec thermal ramp to 94°C, 94°C for 1 min; X 35 cycles: 1 degree/sec thermal ramp to 94°C, then 94°C for 10 sec, followed by 1 degree/sec thermal ramp to 50°C, then 50°C for 10 sec, followed by 1 degree/sec thermal ramp to 60°C, then 60°C for 4 minutes.

The cycle sequencing reaction product was cleaned up to remove the unincorporated dye-labeled terminators that can obscure data at the beginning of the sequence. A precipitation protocol was used. To each sequencing reaction in the 96-well plate, 20 µl of Milli-Q water and 60 µl of 100% isopropanol was added. The plate was left at room temperature for at least 20 minutes to precipitate the extension products. The plate was spun in a plate centrifuge (Jouan) at 3,000 x g for 30 minutes.

Without disturbing the pellet, the supernatant was discarded by inverting the plate onto several paper tissues (Kimwipes) folded to the size of the plate. The inverted plate, with Kimwipes in place, was placed into the centrifuge (Jouan) and spun at 700 x g for 1 minute. The Kimwipes were discarded and the samples were loaded onto a sequencing gel.

Approximately 1 µl of sequencing product was loaded into each well of a 96-lane 5% Long Ranger (FMC single pack) gel. The running buffer consisted of 1X TBE. The glass plates consisted of ABI 48-cm plates for use with a 96-lane 0.4 mm Mylar shark-tooth comb. A semi-automated ABI Prism 377-96 DNA sequencer was used (ABI 377 with 96-lane, Big Dye upgrades). Sequencing run settings were as follows: run module 48E-1200, 8 hr collection time, 2400 V electrophoresis voltage, 50 mA electrophoresis current, 200 W electrophoresis power, CCD offset of 0, gel temperature of 51°C, 40 mW laser power, and CCD gain of 2.

Pyrosequencing is another method of sequencing DNA by synthesis, where the addition of one of the four dNTPs that correctly matches the complementary base on the template strand is detected. Detection occurs via utilization of the pyrophosphate molecules liberated upon base addition to the elongating synthetic strand. The pyrophosphate molecules are used to make ATP, which in turn drives the emission of photons in a luciferin/luciferase reaction, and these photons are detected by the instrument.

A Luc96 Pyrosequencer was used under default operating conditions supplied by the manufacturer. Primers were designed to anneal within 5 bases of the polymorphism, to serve as sequencing primers.

Patient genomic DNA was subject to PCR using amplifying primers that amplify an approximately 200 base pair amplicon containing the polymorphisms of interest. One of the amplifying primers, whose orientation is opposite to the sequencing primer, was biotinylated. This allowed selection of single stranded template for pyrosequencing, whose orientation is complementary to the sequencing primer. Amplicons prepared from genomic DNA were isolated by binding to streptavidin-coated magnetic beads. After denaturation in NaOH, the biotinylated strands were separated from their complementary strands using magnetics. After washing the magnetic beads, the biotinylated template strands still bound to the beads were transferred into 96-well plates. The sequencing primers were added, annealing was carried out at 95° for 2 minutes, and plates were placed in the Pyrosequencer. The enzymes, substrates and dNTPs used for synthesis and pyrophosphate detection were added to the instrument immediately prior to sequencing.

The Luc96 software requires definition of a program of adding the four dNTPs that is specific for the location of the sequencing primer, the DNA composition flanking the SNP, and the two possible alleles at the polymorphic locus. This order of adding the bases generates theoretical outcomes of light intensity patterns for each of the two possible homozygous states and the single heterozygous state. The Luc96 software then compares the actual outcome to the theoretical outcome and calls a genotype for each well. Each sample is also assigned one of three confidence scores: pass, uncertain, fail. The results for each plate are output as a text file and processed in Excel using a Visual Basic program to generate a report of genotype and allele frequencies for the various disease and population cell groupings represented on the 96 well plate.

Prediction of potential transcription binding factor sites was performed using a commercially available software program [GENOMATIX MatInspector Professional; URL: http://genomatix.gsf.de/cgi-bin/matinspector/matinspector.pl ; Quandt et al., *Nucleic Acids Res.*, 23: 4878-4884 (1995)].

Example 2

T→G Transversion at Position 2239 of Human EDN1

Table 1

GROUP I ALLELE FREQUENCY		
	T	**G**
CONTROL		
Black men (n=22 chromosomes)	13 (59%)	9 (41%)
Black women (n=22 chromosomes)	11 (50%)	11 (50%)
White men (n=18 chromosomes)	16 (89%)	2 (11%)
White women (n=24 chromosomes)	21 (88%)	3 (13%)
DISEASE		
BREAST CANCER		
Black women (n=24 chromosomes)	12 (50%)	12 (50%)
White women (n=22 chromosomes)	19 (86%)	3 (14%)
LUNG CANCER		
Black men (n=24 chromosomes)	17 (71%)	7 (29%)
Black women (n=4 chromosomes)	1 (25%)	3 (75%)
White men (n=22 chromosomes)	15 (68%)	7 (32%)
White women (n=20 chromosomes)	14 (70%)	6 (30%)
PROSTATE CANCER		
Black men (n=24 chromosomes)	13 (54%)	11 (46%)
White men (n=24 chromosomes)	17 (71%)	7 (29%)
NIDDM		
Black men (n=20 chromosomes)	14 (70%)	6 (30%)
Black women (n=20 chromosomes)	14 (70%)	6 (30%)
White men (n=22 chromosomes)	16 (73%)	6 (27%)
White women (n=20 chromosomes)	13 (65%)	7 (35%)
ESRD due to NIDDM		
Black men (n=8 chromosomes)	6 (75%)	2 (25%)
Black women (n=20 chromosomes)	14 (70%)	6 (30%)
White men (n=20 chromosomes)	18 (90%)	2 (10%)
White women (n=16 chromosomes)	16 (100%)	0 (0%)

5

Table 2

GROUP II ALLELE FREQUENCY

Disease	Race	CHROMOSOMES	N	T	N	G
Controls	African-American	90	61	67.8%	29	32.2%
	Caucasian	88	76	86.4%	12	13.6%
Colon cancer	African-American	44	31	70.5%	13	29.5%
	Caucasian	44	35	79.5%	9	20.5%
Lung cancer	African-American	40	26	65.0%	14	35.0%
	Caucasian	44	31	70.5%	13	29.5%
Hypertension	African-American	48	31	64.6%	17	35.4%
	Caucasian	44	40	90.9%	4	9.1%
CVA due to HTN	Caucasian	46	38	82.6%	8	17.4%
ESRD due to HTN	African-American	42	26	61.9%	16	38.1%
	Caucasian	48	38	79.2%	10	20.8%
HTN CM	African-American	48	30	62.5%	18	37.5%
	Caucasian	46	38	82.6%	8	17.4%
NIDDM	African-American	42	32	76.2%	10	23.8%
ASPVD due to NIDDM	Caucasian	46	38	82.6%	8	17.4%
CVA due to NIDDM	Caucasian	44	39	88.6%	5	11.4%
ESRD due to NIDDM	Caucasian	46	35	76.1%	11	23.9%
Ischemic CM with NIDDM	African-American	48	30	62.5%	18	37.5%
	Caucasian	48	42	87.5%	6	12.5%
MI due to NIDDM	Caucasian	48	37	77.1%	11	22.9%
Afib without valvular disease	African-American	48	29	60.4%	19	39.6%
	Caucasian	48	40	83.3%	8	16.7%
Alcohol abuse	African-American	48	22	45.8%	26	54.2%
	Caucasian	48	36	75.0%	12	25.0%
Asthma	Caucasian	48	41	85.4%	7	14.6%
COPD	African-American	40	33	82.5%	7	17.5%
	Caucasian	42	34	81.0%	8	19.0%
ESRD due to FSGS	Caucasian	42	33	78.6%	9	21.4%

Table 3

GROUP I GENOTYPE FREQUENCIES			
	T/T	T/G	G/G
CONTROLS			
Black men (n=11)	4 (36%)	5 (45%)	2 (18%)
Black women (n=11)	4 (36%)	3 (27%)	4 (36%)
White men (n=9)	8 (89%)	0 (0%)	1 (11%)
White women (n=12)	9 (75%)	3 (25%)	0 (0%)
DISEASE			
BREAST CANCER			
Black women (n=12)	4 (33%)	4 (33%)	4 (33%)
White women (n=11)	8 (73%)	3 (27%)	0 (0%)
LUNG CANCER			
Black men (n=12)	6 (50%)	5 (42%)	1 (8%)
Black women (n=2)	0 (0%)	1 (50%)	1 (50%)
White men (n=11)	5 (45%)	5 (45%)	1 (9%)
White women (n=10)	5 (50%)	4 (40%)	1 (10%)
PROSTATE CANCER			
Black men (n=12)	3 (25%)	7 (58%)	2 (17%)
White men (n=12)	5 (42%)	7 (58%)	0 (0%)
NIDDM			
Black men (n=10)	6 (60%)	2 (20%)	2 (20%)
Black women (n=10)	5 (50%)	4 (40%)	1 (10%)
White men (n=11)	7 (64%)	2 (18%)	2 (18%)
White women (n=10)	5 (50%)	3 (30%)	2 (20%)
ESRD due to NIDDM			
Black men (n=4)	2 (50%)	2 (50%)	0 (0%)
Black women (n=10)	5 (50%)	4 (40%)	1 (10%)
White men (n=10)	8 (80%)	2 (20%)	0 (0%)
White women (n=8)	8 (100%)	0 (0%)	0 (0%)

Table 4

GROUP II GENOTYPE FREQUENCIES

Disease	Race	People	N	T/T	N	T/G	N	G/G
Controls	African-American	45	17	37.8%	27	60.0%	1	2.2%
	Caucasian	44	33	75.0%	10	22.7%	1	2.3%
Colon cancer	African-American	22	10	45.5%	11	50.0%	1	4.5%
	Caucasian	22	15	68.2%	5	22.7%	2	9.1%
Hypertension	African-American	24	10	41.7%	11	45.8%	3	12.5%
	Caucasian	22	18	81.8%	4	18.2%	0	0.0%
CVA due to HTN	Caucasian	23	16	69.6%	6	26.1%	1	4.3%
ESRD due to HTN	African-American	21	9	42.9%	8	38.1%	4	19.0%
	Caucasian	24	14	58.3%	10	41.7%	0	0.0%
HTN CM	African-American	24	10	41.7%	10	41.7%	4	16.7%
	Caucasian	23	16	69.6%	6	26.1%	1	4.3%
NIDDM	African-American	21	14	66.7%	4	19.0%	3	14.3%
ASPVD due to NIDDM	Caucasian	23	16	69.6%	6	26.1%	1	4.3%
CVA due to NIDDM	Caucasian	22	17	77.3%	5	22.7%	0	0.0%
ESRD due to NIDDM	Caucasian	23	14	60.9%	7	30.4%	2	8.7%
Ischemic CM with NIDDM	African-American	24	10	41.7%	10	41.7%	4	16.7%
	Caucasian	24	18	75.0%	6	25.0%	0	0.0%
MI due to NIDDM	Caucasian	24	13	54.2%	11	45.8%	0	0.0%
Afib without valvular disease	African-American	24	9	37.5%	11	45.8%	4	16.7%
	Caucasian	24	16	66.7%	8	33.3%	0	0.0%
Alcohol abuse	African-American	24	7	29.2%	8	33.3%	9	37.5%
	Caucasian	24	14	58.3%	8	33.3%	2	8.3%
Asthma	Caucasian	24	17	70.8%	7	29.2%	0	0.0%
COPD	African-American	20	13	65.0%	7	35.0%	0	0.0%
	Caucasian	21	14	66.7%	6	28.6%	1	4.8%
ESRD due to FSGS	Caucasian	21	12	57.1%	9	42.9%	0	0.0%

ALLELE-SPECIFIC ODDS RATIOS

5   The susceptibility allele is indicated below, as well as the odds ratio (OR).   The

allele which is present more often in the given disease category was chosen as the

susceptibility allele. For example, the G allele was chosen as the susceptibility allele for

black women with breast cancer because more of the individuals in that category had the

G allele than had the T allele. Where there was a "0" in a cell which produced a 0 in the denominator, Haldane's correction (multiplying all cells by 2 and adding 1) was used. If the odds ratio (OR) was ≥ 1.5, the 95% confidence interval (C.I.) is also given.

An odds ratio of 1.5 was chosen as the threshold of significance based on the recommendation of Austin et al. in *Epidemiol. Rev.*, 16:65-76, 1994. "[E]pidemiology in general and case-control studies in particular are not well suited for detecting weak associations (odds ratios < 1.5)." *Id.* at 66.

An example of the allele-specific odds ratio calculation is given below:

Lung Cancer: Black men

	Cases	Controls
T	17	13
G	7	9

The odds ratio for the T allele is (17)(9)/(7)(13) = 1.7. Therefore, black men with the T allele have a 1.7 fold higher risk of developing lung cancer than black men without the T allele. Odds ratios of 1.5 or higher are highlighted below.

Table 5

GROUP I ALLELE-SPECIFIC ODDS RATIOS			
DISEASE	SUSCEPTIBILITY ALLELE	OR	95% C.I.
Breast Cancer			
Black women	G	1.0	
White women	T	0.9	
Lung Cancer			
Black men	T	*1.7*	0.5-5.7
Black women	G	*3.0*	0.3-33.5
White men	G	*3.7*	0.7-20.9
White women	G	*3.0*	0.6-14.0
Prostate Cancer			
Black men	G	1.3	
White men	G	*3.3*	0.6-18.3
NIDDM			
Black men	T	*1.6*	0.4-5.8
Black women	T	*2.3*	0.7-8.3
White men	G	*3.0*	0.5-17.2
White women	G	*3.8*	0.8-17.2
ESRD due to NIDDM			
Black men	T	1.3*	
Black women	T	1.0*	
White men	T	*3.4**	0.6-19
White women	T	*18.3**	2.3-148

* - as compared to NIDDM.

Table 6
GROUP II ALLELE-SPECIFIC ODDS RATIOS

Disease	Race	Risk Allele	Odds Ratio	Lower Limit 95% CI	Upper Limit 95% CI	Haldane
Colon cancer	Caucasian	C	*1.6*	0.6	4.2	
Hypertension	Caucasian	A	*1.6*	0.5	5.2	
CVA due to HTN*	Caucasian	C	*2.1*	0.6	7.6	
ESRD due to HTN*	African-American	C	1.1	0.5	2.6	
	Caucasian	C	*2.6*	0.8	9.1	
Ischemic CM with NIDDM*[1]	Caucasian	A	*2.1*	0.7	6.2	
Afib without valvular disease	Caucasian	C	1.3	0.5	3.4	
Alcohol abuse	Caucasian	C	*2.1*	0.9	5.2	
Asthma	Caucasian	C	1.1	0.4	3.0	
COPD	Caucasian	C	*1.5*	0.6	4.0	
ESRD due to FSGS	Caucasian	C	*1.7*	0.7	4.5	

GENOTYPE-SPECIFIC ODDS RATIOS

The susceptibility allele (S) is indicated; the alternative allele at this locus is defined as the protective allele (P). Also presented is the odds ratio (OR) for each genotype (SS, SP; the odds ratio for the PP genotype is 1, since it is the reference group, and is not presented separately). For odds ratios $\geq$ 1.5, the 95% confidence interval (C.I.) is also given, in parentheses. Where there was a "0" in a cell which produced a 0 in the denominator, Haldane's correction (multiplying all cells by 2 and adding 1) was used. As discussed above, an odds ratio of 1.5 is chosen as the threshold of significance based on the recommendation of Austin H et al. (Epidemiol. Rev. 16:65-76, 1994).

An example of an odds ratio calculation is worked below, assuming that T is the susceptibility allele (S), and G is the protective allele (P).

Black men: Lung Cancer

	Cases	Controls	Odds Ratio
TT (SS)	6	4	(6)(2)/(1)(4) = 3.0
TG (SP)	5	5	(5)(2)/(1)(5) = 2.0
GG (PP)	1	2	1.0 (by definition)

The odds ratios for individual genotypes are given below. Odds ratios of 1.5 or higher are high-lighted below.

Table 7

GROUP I GENOTYPE-SPECIFIC ODDS RATIOS			
DISEASE	SUSCEPTIBILITY ALLELE	OR(SS)	OR(SP)
Lung Cancer			
Black men	T	_3.0_ (0.2-45.2)	_2.0_ (0.1-29.8)
Black women	G	_3.0_ (0.3-34.6)	_3.9_ (0.3-45.6)
White men	G	_1.5_ (0.3-9.1)	_17.0_ (1.9-151)
White women	G	_5.2_ (0.5-56.1)	_2.2_ (0.6-7.6)
Prostate Cancer			
White men	G	0.5	_23.2_ (2.7-201)
NIDDM			
Black men	T	_1.5_ (0.1-15.5)	0.4
Black women	T	_5.0_ (0.4-64.4)	_5.3_ (0.4-75.8)
White men	G	_1.9_ (0.4-9.3)	_5.7_ (0.6-54.1)
White women	G	_8.6_ (0.9-83.8)	_1.7_ (0.5-6.2)
ESRD due to NIDDM			
White men	T	_5.7_ (0.6-54.1)*	_5.0_ (0.4-59.7)*
White women	T	_7.7_ (0.8-75.3)*	0.7 *

* - as compared to NIDDM

Table 8

GROUP II GENOTYPE-SPECIFIC ODDS RATIOS

Disease	Race	RISK ALLELE	SS O.R.	HALDANE	SP O.R.	HALDANE
Colon cancer	Caucasian	C	0.2		0.3	
Hypertension	Caucasian	A	_1.7_	H	1.3	H
CVA due to HTN*	Caucasian	C	0.0		0.0	
ESRD due to HTN*	African-American	C	0.7		0.5	
	Caucasian	C	0.8	H	_2.3_	H
Ischemic CM with NIDDM*[1]	Caucasian	A	1.4	H	0.6	H
Afib without valvular disease	Caucasian	C	_1.5_	H	_2.4_	H
Alcohol abuse	Caucasian	C	0.2		0.4	
Asthma	Caucasian	C	_1.6_	H	_2.1_	H
COPD	Caucasian	C	0.4		0.6	
ESRD due to FSGS	Caucasian	C	1.1	H	_2.7_	H

*-Compared to HTN alone.
*[1]-Compared to MI with NIDDM.

PCR and sequencing were conducted as described in Example 1. The primers used were the same as in Example 1. The control samples are in good agreement with Hardy-Weinberg equilibrium, as follows:

5   For the Group I diseases, a frequency of 0.59 for the T allele ("p") and 0.41 for the G allele ("q") among black male control individuals predicts genotype frequencies of 35% T/T, 48% T/G, and 17% G/G at Hardy-Weinberg equilibrium ($p^2 + 2pq + q^2 = 1$). The observed genotype frequencies were 36% T/T, 45% T/G, and 18% C/C, in close agreement with those predicted for Hardy-Weinberg equilibrium.

10   A frequency of 0.50 for the T allele ("p") and 0.50 for the G allele ("q") among black female control individuals predicts genotype frequencies of 25% T/T, 50% T/G, and 25% G/G at Hardy-Weinberg equilibrium ($p^2 + 2pq + q^2 = 1$). The observed genotype frequencies were 36% T/T, 27% T/G, and 36% C/C, in rather distant agreement with those predicted for Hardy-Weinberg equilibrium.

15   A frequency of 0.89 for the T allele ("p") and 0.11 for the G allele ("q") among white male control individuals predicts genotype frequencies of 79% T/T, 20% T/G, and 1% G/G at Hardy-Weinberg equilibrium ($p^2 + 2pq + q^2 = 1$). The observed genotype frequencies were 89% T/T, 0% T/G, and 11% C/C, in rather distant agreement with those predicted for Hardy-Weinberg equilibrium.

20   A frequency of 0.88 for the T allele ("p") and 0.13 for the G allele ("q") among white female control individuals predicts genotype frequencies of 77% T/T, 21% T/G, and 2% G/G at Hardy-Weinberg equilibrium ($p^2 + 2pq + q^2 = 1$). The observed genotype frequencies were 75% T/T, 25% T/G, and 0% C/C, in close agreement with those predicted for Hardy-Weinberg equilibrium.

25   For the Group II diseases, a frequency of 0.68 for the T allele ("p") and 0.32 for the G allele ("q") among African American control individuals predicts genotype frequencies of 45.9% T/T, 44.0% T/G, and 10.1% G/G at Hardy-Weinberg equilibrium ($p^2 + 2pq + q^2 = 1$). The observed genotype frequencies were 37.8% T/T, 60.0% T/G, and 2.2% G/G, in distant agreement with those predicted for Hardy-Weinberg equilibrium.

30   A frequency of 0.86 for the T allele ("p") and 0.14 for the G allele ("q") among Caucasian control individuals predicts genotype frequencies of 74.6% T/T, 23.5% T/G, and 1.9% G/G at Hardy-Weinberg equilibrium ($p^2 + 2pq + q^2 = 1$). The observed genotype frequencies were 75.0% T/T, 22.7% T/G, and 2.3% G/G, in excellent agreement with those predicted for Hardy-Weinberg equilibrium.

Using an allele-specific odds ratio of 1.5 or greater as a practical level of significance (see Austin H et al., discussed above), the following observations can be made.

For black men with lung cancer, the odds ratio for the T allele as a risk factor for disease is 1.7 (95% CI, 0.5-5.7). The odds ratio for the homozygote (TT) is 3.0 (95% CI, 0.2- 45.2). The heterozygote (TG genotype) has an odds ratio of 2.0 (95% C.I., 0.1-29.8). These data suggest that the T allele behaves as a dominant allele, with an additive effect of allele dosage (2.0 + 2.0 - 1 = 3.0).

For black women with lung cancer, the odds ratio for the G allele as a risk factor for disease is 3.0 (95% CI, 0.3-33.5). The odds ratio for the homozygote (GG) is 3.0 (95% CI, 0.3-34.6). The heterozygote (GT genotype) has an odds ratio of 3.9 (95% C.I., 0.3-45.6). T hese data suggest that the G allele behaves as a dominant allele, with no additional effect of having two copies of the G allele (GG homozygote) as compared with having only one copy (GT heterozygote).

For white men with lung cancer, the odds ratio for the G allele as a risk factor for disease is 3.7 (95% CI, 0.7-20.9). The odds ratio for the homozygote (GG) is only 1.5 (95% CI, 0.3-9.1), whereas the heterozygote (GT genotype) has a remarkable odds ratio of 17.0 (95% C.I., 1.9-151). These data suggest that the G allele behaves as a codominant allele.

For white women with lung cancer, the odds ratio for the G allele as a risk factor for disease is 3.0 (95% CI, 0.6-14.0). The odds ratio for the homozygote (GG) is 5.2 (95% CI, 0.5-56.1), while the heterozygote (GT genotype) has an odds ratio of 2.2 (95% C.I., 0.6-7.6). These data suggest that the G allele behaves as a dominant allele with more than an additive effect of allele copy number (2.2 + 2.2 - 1 < 5.2).

For white men with prostate cancer, the odds ratio for the G allele as a risk factor for disease is 3.3 (95% CI, 0.6-18.3). The odds ratio for the homozygote (GG) is actually less than 1, whereas the heterozygote (GT genotype) has a remarkable odds ratio of 23.2 (95% C.I., 2.7-201). These data suggest that the G allele behaves as a codominant allele.

For black men with NIDDM, the odds ratio for the T allele as a risk factor for disease is 1.6 (95% CI, 0.4-5.8). The odds ratio for the homozygote (TT) is 1.5 (95% CI, 0.1-15.5), whereas the heterozygote (TG genotype) has an odds ratio of less than 1. These data suggest that the T allele behaves as a recessive allele.

For black women with NIDDM, the odds ratio for the T allele as a risk factor for disease is 2.3 (95% CI, 0.7-8.3). The odds ratio for the homozygote (TT) is 5.0 (95% CI,

0.4-64.4), whereas the heterozygote (TG genotype) has an odds ratio of 5.3 (95% CI, 0.4-75.8). These data suggest that the T allele behaves as a classical dominant allele.

For white men with NIDDM, the odds ratio for the G allele as a risk factor for disease is 3.0 (95% CI, 0.5-17.2). The odds ratio for the homozygote (GG) is 1.9 (95% CI, 0.4-9.3), whereas the heterozygote (GT genotype) has an odds ratio of 5.7 (95% CI, 0.6-54.1). These data suggest that the G allele behaves as a codominant allele.

For white women with NIDDM, the odds ratio for the G allele as a risk factor for disease is 3.8 (95% CI, 0.8-17.2). The odds ratio for the homozygote (GG) is 8.6 (95% CI, 0.9-83.8), whereas the heterozygote (GT genotype) has an odds ratio of only 1.7 (95% CI, 0.5-6.2). These data suggest that the G allele behaves as a dominant allele, with a more than multiplicative effect of allele dosage [8.6 >> (1.7)(1.7)].

For white men with ESRD due to NIDDM, the odds ratio for the T allele as a risk factor for disease is 3.4 (95% CI, 0.6-19.2) as compared with white men with NIDDM but no renal disease. The odds ratio for the homozygote (TT) is 5.7 (95% CI, 0.6-54.1), while the heterozygote (TG genotype) has a similar odds ratio of 5.0 (95% CI, 0.4-59.7). These data suggest that the T allele behaves as a classical dominant allele.

For white women with ESRD due to NIDDM, the odds ratio for the T allele as a risk factor for disease is a remarkable 18.3 (95% CI, 2.3-148) as compared with white women with NIDDM but no renal disease. The odds ratio for the homozygote (TT) is 7.7 (95% CI, 0.8-75.3), while the heterozygote (TG genotype) has an odds ratio of only 0.7. These data suggest that the T allele behaves as a classical recessive allele.

For Caucasians with alcohol abuse the odds ratio for the G allele was 2.1 (95% CI, 0.9 - 5.2). Data were not sufficient to generate genotypic odds ratios of 1.5 or greater. These data further suggest that the EDN-1 gene is significantly associated with alcohol abuse in Caucasians, i.e. abnormal activity of the EDN-1 gene predisposes Caucasians to alcohol abuse.

For Caucasians with colon cancer the odds ratio for the G allele was 1.6 (95 % CI, 0.6 - 4.2). Data were not sufficient to generate genotypic odds ratios of 1.5 or greater. These data further suggest that the EDN-1 gene is significantly associated with colon cancer in Caucasians, i.e. abnormal activity of the EDN-1 gene predisposes Caucasians to colon cancer.

For Caucasians with diabetic cardiomyopathy the odds ratio for the T allele was 2.1 (95% CI, 0.7 - 6.2), compared to Caucasians with MI due to NIDDM. Data were not sufficient to generate genotypic odds ratios of 1.5 or greater. These data further suggest

that the EDN-1 gene is significantly associated with diabetic cardiomyopathy in Caucasians, i.e. abnormal activity of the EDN-1 gene predisposes Caucasians to diabetic cardiomyopathy.

For Caucasians with ESRD due to hypertension the odds ratio for the G allele was 2.6 (95% CI, 0.8 - 9.1), compared to Caucasians with hypertension only. The odds ratio for the homozygote (G/G) was 0.8^H (95% CI, 0 - 14.1), while the odds ratio for the heterozygote (T/G) was 2.3^H (95% CI, 0 - 137). These data suggest that G allele acts in a co-dominant manner in this patient population. These data further suggest that the EDN-1 gene is significantly associated with ESRD due to hypertension in Caucasians, i.e. abnormal activity of the EDN-1 gene predisposes Caucasians to ESRD due to hypertension.

For Caucasians with ESRD due to FSGS the odds ratio for the G allele was 1.7 (95% CI, 0.7 - 4.5). The odds ratio for the homozygote (G/G) was 1.1^H (95% CI, 0.1 - 19.7), while the odds ratio for the heterozygote (T/G) was 2.7^H (95% CI, 0.1 - 75). These data suggest that the G allele acts in a co-dominant manner in this patient population. These data further suggest that the EDN-1 gene is significantly associated with ESRD due to FSGS in Caucasians, i.e. abnormal activity of the EDN-1 gene predisposes Caucasians to ESRD due to FSGS.

For Caucasians with hypertension only the odds ratio for the T allele was 1.6 (95% CI, 0.5 - 5.2). The odds ratio for the homozygote (T/T) was 1.7^H (95% CI, 0.1 - 28.6), while the odds ratio for the heterozygote (T/G) was 1.3^H (95% CI, 0 - 38). These data suggest that the T allele acts in a recessive manner in this patient population. These data further suggest that the EDN-1 gene is significantly associated with hypertension only in Caucasians, i.e. abnormal activity of the EDN-1 gene predisposes Caucasians to hypertension only.

For Caucasians with CVA due to HTN the odds ratio for the G allele was 2.1 (95% CI, 0.6 - 7.6), compared to Caucasians with hypertension only. Data were not sufficient to generate genotypic odds ratios of 1.5 or greater. These data further suggest that the EDN-1 gene is significantly associated with CVA due to HTN in Caucasians, i.e. abnormal activity of the EDN-1 gene predisposes Caucasians to CVA due to HTN.

The binding site of T-cell factor-2 alpha (TCF-2 alpha) is predicted to be disrupted by the T2239→G SNP (Quandt K et al., *Nucleic Acids Res.*, 23:4878-4884, 1995). TCF-2 alpha binds to a core sequence of five nucleotides, 5'-KTKTC-3' (Waterman ML, et al. *New Biology*, 2(7):621-636, 1990). A TCF-2 alpha binding site, which occurs on average

3.91 times per 1000 base pairs of random genomic sequence in vertebrates, is predicted to occur at position 2236 to 2240 on the (-) strand of reference sequence J05008.1 (matrix score 1.000, with 1.000 being an identical match). The T2239-->G SNP replaces the indicated _T_ with a G within the core binding sequence.

5   TCF-2 alpha is a transcriptional activator in lymphoid cells, although nothing is known of its activity in other cell types. Disruption of the TCF-2 alpha core binding site is expected to result in a decreased rate of transcription of the endothelin-1 gene.

Example 3

A to C Substitution at Position 2657 of Human EDN-1 Promoter

Table 9

GROUP I ALLELE FREQUENCIES		
CONTROL	A	C
Black men (n=46 chromosomes)	30 (65%)	16 (35%)
Black women (n=40 chromosomes)	30 (75%)	10 (25%)
White men (n=42 chromosomes)	38 (90%)	4 (10%)
White women (n=48 chromosomes)	42 (88%)	6 (13%)
DISEASE		
Breast Cancer		
Black women (n=16 chromosomes)	7 (44%)	9 (56%)
White women (n=12 chromosomes)	9 (75%)	3 (25%)
Lung Cancer		
Black men (n=20 chromosomes)	13 (65%)	7 (35%)
Black women (n=16 chromosomes)	11 (69%)	5 (31%)
White men (n=20 chromosomes)	13 (65%)	7 (35%)
White women (n=12 chromosomes)	8 (67%)	4 (33%)
Prostate Cancer		
Black men (n=16 chromosomes)	13 (81%)	3 (19%)
White men (n=18 chromosomes)	11 (61%)	7 (39%)
HTN		
Black men (n=18 chromosomes)	12 (67%)	6 (33%)
Black women (n=16 chromosomes)	13 (81%)	3 (19%)
White men (n=22 chromosomes)	21 (95%)	1 (5%)
White women (n=18 chromosomes)	15 (83%)	3 (17%)
ESRD due to HTN		
Black men (n=12 chromosomes)	10 (83%)	2 (17%)
Black women (n=10 chromosomes)	6 (60%)	4 (40%)
White men (n=14 chromosomes)	12 (86%)	2 (14%)
White women (n=4 chromosomes)	4 (100%)	0 (0%)
NIDDM		
Black men (n=16 chromosomes)	13 (81%)	3 (19%)
Black women (n=16 chromosomes)	11 (69%)	5 (31%)
White men (n=22 chromosomes)	16 (73%)	6 (27%)
White women (n=20 chromosomes)	15 (75%)	5 (25%)
ESRD due to NIDDM		
Black men (n=4 chromosomes)	3 (75%)	1 (25%)
Black women (n=18 chromosomes)	14 (78%)	4 (22%)
White men (n=16 chromosomes)	14 (88%)	2 (13%)
White women (n=10 chromosomes)	10 (100%)	0 (0%)

Table 10

GROUP II ALLELE FREQUENCIES

Disease	Race	CHROMOSOMES	N	C	N	A
Controls	African-American	90	25	27.8%	65	72.2%
	Caucasian	90	15	16.7%	75	83.3%
Colon cancer	African-American	48	8	16.7%	40	83.3%
	Caucasian	44	7	15.9%	37	84.1%
Hypertension	African-American	42	6	14.3%	36	85.7%
	Caucasian	44	4	9.1%	40	90.9%
ASPVD due to HTN	African-American	50	10	20.0%	40	80.0%
	Caucasian	50	7	14.0%	43	86.0%
CVA due to HTN	Caucasian	48	9	18.8%	39	81.3%
Cataracts due to HTN	African-American	44	9	20.5%	35	79.5%
HTN CM	African-American	48	7	14.6%	41	85.4%
	Caucasian	44	5	11.4%	39	88.6%
MI due to HTN	African-American	42	11	26.2%	31	73.8%
	Caucasian	46	11	23.9%	35	76.1%
NIDDM	African-American	44	11	25.0%	33	75.0%
	Caucasian	48	13	27.1%	35	72.9%
ASPVD due to NIDDM	African-American	46	15	32.6%	31	67.4%
	Caucasian	46	8	17.4%	38	82.6%
CVA due to NIDDM	African-American	48	9	18.8%	39	81.3%
	Caucasian	46	5	10.9%	41	89.1%
Ischemic CM	African-American	48	11	22.9%	37	77.1%
	Caucasian	42	8	19.0%	34	81.0%
Ischemic CM with NIDDM	African-American	48	14	29.2%	34	70.8%
	Caucasian	48	7	14.6%	41	85.4%
MI due to NIDDM	African-American	48	6	12.5%	42	87.5%
	Caucasian	46	10	21.7%	36	78.3%
Afib without valvular disease	African-American	48	14	29.2%	34	70.8%
	Caucasian	48	8	16.7%	40	83.3%
Alcohol abuse	African-American	48	17	35.4%	31	64.6%
	Caucasian	48	12	25.0%	36	75.0%

	CHROMOSOMES	N	C	N	A
Anxiety	African-American	48	16	33.3%	32	66.7%
	Caucasian	42	10	23.8%	32	76.2%
Asthma	African-American	48	11	22.9%	37	77.1%
	Caucasian	48	6	12.5%	42	87.5%
COPD	African-American	44	3	6.8%	41	93.2%
	Caucasian	42	8	19.0%	34	81.0%
Cholecystectomy	African-American	48	14	29.2%	34	70.8%
	Caucasian	46	7	15.2%	39	84.8%
DJD	African-American	40	9	22.5%	31	77.5%
ESRD and frequent de-clots	African-American	46	13	28.3%	33	71.7%
	Caucasian	42	5	11.9%	37	88.1%
ESRD due to FSGS	African-American	44	13	29.5%	31	70.5%
	Caucasian	46	10	21.7%	36	78.3%
ESRD due to IDDM	African-American	48	14	29.2%	34	70.8%
	Caucasian	44	3	6.8%	41	93.2%
Seizure disorder	African-American	46	19	41.3%	27	58.7%
	Caucasian	48	5	10.4%	43	89.6%

Table 11

GROUP I GENOTYPE FREQUENCIES			
	A/A	A/C	C/C
CONTROLS			
Black men (n=23)	10 (43%)	10 (43%)	3 (13%)
Black women (n=20)	11 (55%)	8 (40%)	1 (5%)
White men (n=21)	18 (86%)	2 (10%)	1 (5%)
White women (n=24)	19 (79%)	4 (17%)	1 (4%)
DISEASE			
Breast Cancer			
Black women (n=8)	3 (38%)	1 (13%)	4 (50%)
White women (n=6)	4 (67%)	1 (17%)	1 (17%)
Lung Cancer			
Black men (n=10)	5 (50%)	3 (30%)	2 (20%)
Black women (n=8)	4 (50%)	3 (38%)	1 (13%)
White men (n=10)	5 (50%)	3 (30%)	2 (20%)
White women (n=6)	2 (33%)	4 (67%)	0 (0%)
Prostate Cancer			
Black men (n=8)	5 (63%)	3 (38%)	0 (0%)
White men (n=9)	3 (33%)	5 (56%)	1(11%)
HTN			
Black men (n=9)	5 (56%)	2 (22%)	2 (22%)
Black women (n=8)	5 (63%)	3 (38%)	0 (0%)
White men (n=11)	10 (91%)	1 (9%)	0 (0%)
White women (n=9)	7 (78%)	1 (11%)	1 (11%)
ESRD due to HTN			
Black men (n=6)	4 (67%)	2 (33%)	0 (0%)
Black women (n=5)	3 (60%)	0 (0%)	2 (40%)
White men (n=7)	5 (71%)	2 (29%)	0 (0%)
White women (n=2)	2 (100%)	0 (0%)	0 (0%)
NIDDM			
Black men (n=8)	5 (63%)	3 (38%)	0 (0%)
Black women (n=8)	4 (50%)	3 (38%)	1 (13%)
White men (n=11)	7 (64%)	2 (18%)	2 (18%)
White women (n=10)	6 (60%)	3 (30%)	1 (10%)
ESRD due to NIDDM			
Black men (n=2)	1 (50%)	1 (50%)	0 (0%)
Black women (n=9)	5 (56%)	4 (44%)	0 (0%)
White men (n=8)	6 (75%)	2 (25%)	0 (0%)
White women (n=5)	5 (100%)	0 (0%)	0 (0%)

Table 12

GROUP II GENOTYPE FREQUENCIES

Disease	Race	People	N	C/C	N	C/A	N	A/A
Controls	African-American	45	4	8.9%	17	37.8%	24	53.3%
	Caucasian	45	3	6.7%	9	20.0%	33	73.3%
Colon cancer	African-American	24	0	0.0%	8	33.3%	16	66.7%
	Caucasian	22	1	4.5%	5	22.7%	16	72.7%
Hypertension	African-American	21	0	0.0%	6	28.6%	15	71.4%
	Caucasian	22	0	0.0%	4	18.2%	18	81.8%
ASPVD due to HTN	African-American	25	2	8.0%	6	24.0%	17	68.0%
	Caucasian	25	1	4.0%	5	20.0%	19	76.0%
CVA due to HTN	Caucasian	24	1	4.2%	7	29.2%	16	66.7%
Cataracts due to HTN	African-American	22	1	4.5%	7	31.8%	14	63.6%
ESRD due to HTN	African-American	22	4	18.2%	8	36.4%	10	45.5%
	Caucasian	24	1	4.2%	10	41.7%	13	54.2%
HTN CM	African-American	24	2	8.3%	3	12.5%	19	79.2%
	Caucasian	22	0	0.0%	5	22.7%	17	77.3%
MI due to HTN	African-American	21	2	9.5%	7	33.3%	12	57.1%
	Caucasian	23	2	8.7%	7	30.4%	14	60.9%
NIDDM	African-American	22	2	9.1%	7	31.8%	13	59.1%
	Caucasian	24	5	20.8%	3	12.5%	16	66.7%
ASPVD due to NIDDM	African-American	23	2	8.7%	11	47.8%	10	43.5%
	Caucasian	23	1	4.3%	6	26.1%	16	69.6%
CVA due to NIDDM	African-American	24	0	0.0%	9	37.5%	15	62.5%
	Caucasian	23	0	0.0%	5	21.7%	18	78.3%
Ischemic CM	African-American	24	2	8.3%	7	29.2%	15	62.5%
	Caucasian	21	1	4.8%	6	28.6%	14	66.7%
Ischemic CM with NIDDM	African-American	24	3	12.5%	8	33.3%	13	54.2%
	Caucasian	24	0	0.0%	7	29.2%	17	70.8%
MI due to NIDDM	African-American	24	0	0.0%	6	25.0%	18	75.0%
	Caucasian	23	0	0.0%	10	43.5%	13	56.5%
Afib without valvular disease	African-American	24	1	4.2%	12	50.0%	11	45.8%
	Caucasian	24	0	0.0%	8	33.3%	16	66.7%
Alcohol abuse	African-American	24	5	20.8%	7	29.2%	12	50.0%
	Caucasian	24	2	8.3%	8	33.3%	14	58.3%

	People	N	C/C	N	C/A	N	A/A
Anxiety	African-American	24	3	12.5%	10	41.7%	11	45.8%
	Caucasian	21	0	0.0%	10	47.6%	11	52.4%
Asthma	African-American	24	2	8.3%	7	29.2%	15	62.5%
	Caucasian	24	0	0.0%	6	25.0%	18	75.0%
COPD	African-American	22	0	0.0%	3	13.6%	19	86.4%
	Caucasian	21	1	4.8%	6	28.6%	14	66.7%
Cholecystectomy	African-American	24	1	4.2%	12	50.0%	11	45.8%
	Caucasian	23	0	0.0%	7	30.4%	16	69.6%
DJD	African-American	20	1	5.0%	7	35.0%	12	60.0%
ESRD and frequent de-clots	African-American	23	3	13.0%	7	30.4%	13	56.5%
	Caucasian	21	1	4.8%	3	14.3%	17	81.0%
ESRD due to FSGS	African-American	22	1	4.5%	11	50.0%	10	45.5%
	Caucasian	23	0	0.0%	10	43.5%	13	56.5%
ESRD due to IDDM	African-American	24	3	12.5%	8	33.3%	13	54.2%
	Caucasian	22	0	0.0%	3	13.6%	19	86.4%
Seizure disorder	African-American	23	4	17.4%	11	47.8%	8	34.8%
	Caucasian	24	0	0.0%	5	20.8%	19	79.2%

Allele-Specific Odds Ratios

The susceptibility allele is indicated below, as well as the odds ratio (OR). Where there was a "0" in a cell which produced a 0 in the denominator, Haldane's correction (multiplying all cells by 2 and adding 1) was used. If the odds ratio (OR) was ≥ 1.5, the 95% confidence interval (C.I.) is also given. An odds ratio of 1.5 was chosen as the threshold of significance based on the recommendation of Austin et al. in *Epidemiol. Rev.*, 16:65-76, 1994. "[E]pidemiology in general and case-control studies in particular are not well suited for detecting weak associations (odds ratios < 1.5)." *Id.* at 66. Odds ratios of greater than 1.5 are highlighted below.

Table 13

GROUP I ALLELE-SPECIFIC ODDS RATIOS			
DISEASE	SUSCEPTIBILITY ALLELE	OR	95% C.I.
Breast Cancer			
Black women	C	*3.9*	1.1-13
White women	C	*2.3*	0.5-11
Lung Cancer			
Black men	C	1.0	0.3-3.0
Black women	C	1.4	0.4-4.9
White men	C	*5.1*	1.3-20
White women	C	*3.5*	0.8-15
Prostate Cancer			
Black men	A	*2.3*	0.6-9.3
White men	C	*6.0*	1.5-25
Hypertension (HTN)			
Black men	A	1.1	0.3-3.4
Black women	A	1.4	0.3-6.1
White men	A	*2.2*	0.2-21
White women	C	1.4	0.3-6.3
ESRD due to HTN*			
Black men	A	*2.5*	0.4-15.2
Black women	C	*2.9*	0.5-17.2
White men	C	*3.5*	0.3-42.8
White women	A	*2.0*[H]	0.2-18.8
NIDDM			
Black men	A	*2.3*	0.6-9.3
Black women	C	1.4	
White men	C	*3.6*	0.9-14.4
White women	C	*2.3*	0.6-8.8
ESRD due to NIDDM*[I]			
Black men	C	1.4	
Black women	A	*1.6*	0.3-7.4
White men	A	*2.6*	0.5-15.2
White women	A	*7.5*[H]	0.9-62.1

* - Compared to HTN alone.

*[I] - Compared to NIDDM alone.

Table 14
GROUP II ALLELE-SPECIFIC ODDS RATIOS

Disease	Race	Risk Allele	Odds Ratio	Lower Limit 95% CI	Upper Limit 95% CI	Haldane
Colon cancer	African-American	A	*1.9*	0.8	4.7	
	Caucasian	A	1.1	0.4	2.8	
ASPVD due to HTN*	African-American	C	*1.5*	0.5	4.5	
	Caucasian	C	*1.6*	0.4	6.0	
CVA due to HTN*	Caucasian	C	*2.3*	0.7	8.1	
Cataracts due to HTN*	African-American	A	*1.5*	0.6	3.6	
HTN CM*[1]	African-American	A	*2.1*	0.7	6.0	
	Caucasian	A	*2.5*	0.8	7.8	
MI due to HTN*	African-American	C	*2.1*	0.7	6.4	
	Caucasian	C	*3.1*	0.9	10.8	
ASPVD due to NIDDM*[2]	African-American	C	*1.5*	0.6	3.6	
	Caucasian	A	*1.8*	0.7	4.8	
CVA due to NIDDM*[2]	African-American	A	1.4	0.5	3.9	
	Caucasian	A	*3.0*	1.0	9.4	
Ischemic CM with NIDDM*[3]	African-American	C	*2.9*	1.0	8.3	
	Caucasian	A	*1.6*	0.6	4.7	
MI due to NIDDM*[2]	African-American	A	*2.3*	0.8	7.0	
	Caucasian	A	1.3	0.5	3.4	
Afib without valvular disease	African-American	C	1.1	0.5	2.3	
	Caucasian	A	1.0	0.4	2.6	
Alcohol abuse	African-American	C	1.4	0.7	3.0	
	Caucasian	C	*1.7*	0.7	3.9	
Anxiety	African-American	C	1.3	0.6	2.8	
	Caucasian	C	*1.6*	0.6	3.8	
Asthma	African-American	A	1.3	0.6	2.9	
	Caucasian	A	1.4	0.5	3.9	
COPD	African-American	A	*5.3*	1.5	18.5	
	Caucasian	C	1.2	0.5	3.0	
Cholecystectomy	African-American	C	1.1	0.5	2.3	
	Caucasian	A	1.1	0.4	3.0	

		Risk Allele	Odds Ratio	Lower Limit 95% CI	Upper Limit 95% CI	Haldane
DJD	African-American	A	1.3	0.6	3.2	
ESRD and frequent de-clots	African-American	C	1.0	0.5	2.3	
	Caucasian	A	_1.5_	0.5	4.4	
ESRD due to FSGS	African-American	C	1.1	0.5	2.4	
	Caucasian	C	1.4	0.6	3.4	
ESRD due to IDDM	African-American	C	1.1	0.5	2.3	
	Caucasian	A	_2.7_	0.7	10.0	
Seizure disorder	African-American	C	_1.8_	0.9	3.9	
	Caucasian	A	_1.7_	0.6	5.1	

*-Compared to HTN alone.
*[1]-Compared to MI with HTN.
*[2]-Compared to NIDDM alone.
*[3]-Compared to MI with NIDDM.

Genotype-Specific Odds Ratios

The susceptibility allele (S) is indicated; the alternative allele at this locus is defined as the protective allele (P). Also presented is the odds ratio (OR) for each genotype (SS, SP). The odds ratio for the PP genotype is 1, since it is the reference group, and is not presented separately. For odds ratios ≥ 1.5, the 95% confidence interval (C.I.) is also given, in parentheses. An odds ratio of 1.5 was chosen as the threshold of significance based on the recommendation of Austin et al., in *Epidemiol. Rev.*, 16:65-76, 1994. "[E]pidemiology in general and case-control studies in particular are not well suited for detecting weak associations (odds ratios < 1.5)." *Id.* at 66.

Haldane's zero cell correction was used when the denominator contained a zero. Odds ratios of greater than 1.5 are highlighted below.

Table 15

GROUP I GENOTYPE-SPECIFIC ODDS RATIOS			
DISEASE	SUSCEPTIBILITY ALLELE	OR(SS)	OR(SP)
Breast Cancer			
Black women	C	_14.7_ (1.2-185)	0.5 (0-5.3)
White women	C	_4.8_ (0.2-93)	1.2 (0.1-14)
Lung Cancer			
White men	C	_7.2_ (0.5-97)	_5.4_ (0.7-42)
White women	C	0	_9.5_ (1.3-71.0)
Prostate Cancer			
Black men	A	_3.7_ (0.4-34)[H]	_2.3_ (0.2-22)[H]
White men	C	_6.0_ (0.3-124)	_15.0_ (1.9-116)
Hypertension (HTN)			
White men	A	_1.7_ (0.2-17)[H]	_1.8_ (0.1-26)[H]
ESRD due to HTN *			
Black men	A	_4.1_ (0.4-42)[H]	_5.0_ (0.4-60)[H]
Black women	C	_7.9_ (0.8-82)[H]	0
White men	C	_1.9_ (0.1-34)[H]	_4.0_ (0.3-55.5)[H]
White women	A	1.0[H]	1.0[H]
NIDDM			
Black men	A	_3.7_ (0.4-34)[H]	_2.3_ (0.2-22)[H]
White men	C	_5.1_ (0.4-66)	_2.6_ (0.3-22)
White women	C	_3.2_ (0.2-59)	_2.4_ (0.4-14)
ESRD due to NIDDM *[1]			
Black women	A	_3.7_ (0.3-42)[H]	_3.9_ (0.3-46)[H]
White men	A	_4.3_ (0.4-42)[H]	_5.0_ (0.4-60)[H]
White women	A	_2.5_ (0.2-28)[H]	0.4 (0-9.4)[H]

Table 16

GROUP II GENOTYPE-SPECIFIC ODDS RATIOS

Disease	Race	RISK ALLELE	SS O.R.	HALDANE	SP O.R.	HALDANE
Colon cancer	African-American	A	0.0		0.7	
	Caucasian	A	0.7		1.1	
ASPVD due to HTN*	African-American	C	4.4	H	0.9	
	Caucasian	C	2.8	H	1.2	
CVA due to HTN*	Caucasian	C	3.4	H	2.0	
Cataracts due to HTN*	African-American	A	0.4		0.7	
HTN CM*[1]	African-American	A	0.6		0.3	
	Caucasian	A	0.0		0.6	
MI due to HTN*	African-American	C	6.2	H	1.5	
	Caucasian	C	6.4	H	2.3	
ASPVD due to NIDDM*[2]	African-American	C	1.3		2.0	
	Caucasian	A	0.2		2.0	
CVA due to NIDDM*[2]	African-American	A	0.0		1.1	
	Caucasian	A	0.0		1.5	
Ischemic CM with NIDDM*[3]	African-American	C	9.6	H	1.8	
	Caucasian	A	0.8	H	0.5	
MI due to NIDDM*[2]	African-American	A	0.0		0.6	
	Caucasian	A	0.0		4.1	
Afib without valvular disease	African-American	C	0.5		1.5	
	Caucasian	A	0.0		1.8	
Alcohol abuse	African-American	C	2.5		0.8	
	Caucasian	C	1.6		2.1	
Anxiety	African-American	C	1.6		1.3	
	Caucasian	C	0.0		3.3	
Asthma	African-American	A	0.8		0.7	
	Caucasian	A	0.0		1.2	
COPD	African-American	A	0.0		0.2	
	Caucasian	C	0.8		1.6	
Cholecystectomy	African-American	C	0.5		1.5	
	Caucasian	A	0.0		1.6	
DJD	African-American	A	0.5		0.8	

		RISK ALLELE	SS O.R.	HALDANE	SP O.R.	HALDANE
ESRD and frequent de-clots	African-American	C	1.4		0.8	
	Caucasian	A	0.6		0.6	
ESRD due to FSGS	African-American	C	0.6		1.6	
	Caucasian	C	0.0		2.8	
ESRD due to IDDM	African-American	C	1.4		0.9	
	Caucasian	A	0.0		0.6	
Seizure disorder	African-American	C	3.0		1.9	
	Caucasian	A	0.0		1.0	

*-Compared to HTN alone.
*[1]-Compared to MI with HTN.
*[2]-Compared to NIDDM alone.
*[3]-Compared to MI with NIDDM.

PCR and sequencing were conducted as described in Example 1. The primers used were those in Example 1. The control samples were in good agreement with Hardy-Weinberg equilibrium, as follows:

For the Group I diseases, a frequency of 0.65 for the A allele ("p") and 0.35 for the C allele ("q") among black male control individuals predicts genotype frequencies of 42% A/A, 46% A/C, and 12% C/C at Hardy-Weinberg equilibrium ($p2 + 2pq + q2 = 1$). The observed genotype frequencies were 43% A/A, 43% A/C, and 13% C/C, in close agreement with those predicted for Hardy-Weinberg equilibrium.

A frequency of 0.75 for the A allele ("p") and 0.25 for the C allele ("q") among black female control individuals predicts genotype frequencies of 56% A/A, 38% A/C, and 6% C/C at Hardy-Weinberg equilibrium ($p2 + 2pq + q2 = 1$). The observed genotype frequencies were 55% A/A, 40% A/C, and 5% C/C, in very close agreement with those predicted for Hardy-Weinberg equilibrium.

A frequency of 0.90 for the A allele ("p") and 0.10 for the C allele ("q") among white male control individuals predicts genotype frequencies of 81% A/A, 18% A/C, and 1% C/C at Hardy-Weinberg equilibrium ($p2 + 2pq + q2 = 1$). The observed genotype frequencies were 86% A/A, 10% A/C, and 5% C/C, in reasonably close agreement with those predicted for Hardy-Weinberg equilibrium.

A frequency of 0.88 for the A allele ("p") and 0.13 for the C allele ("q") among white female control individuals predicts genotype frequencies of 77% A/A, 21% A/C,

and 2% C/C at Hardy-Weinberg equilibrium (p2 + 2pq + q2 = 1). The observed genotype frequencies were 79% A/A, 17% A/C, and 4% C/C, in reasonably close agreement with those predicted for Hardy-Weinberg equilibrium.

For the Group II diseases, a frequency of 0.28 for the C allele ("p") and 0.72 for the A allele ("q") among African-American control individuals predicts genotype frequencies of 8.0% C/C, 40.0% C/A, and 52.0% A/A at Hardy-Weinberg equilibrium (p^2 + 2pq + q^2 = 1). The observed genotype frequencies were 8.9% C/C, 37.8% C/A, and 53.3% A/A, in excellent agreement with those predicted for Hardy-Weinberg equilibrium.

A frequency of 0.17 for the C allele ("p") and 0.83 for the A allele ("q") among Caucasian control individuals predicts genotype frequencies of 3.0% C/C, 28.0% C/A, and 69.0% A/A at Hardy-Weinberg equilibrium (p^2 + 2pq + q^2 = 1). The observed genotype frequencies were 6.7% C/C, 20.0% C/A, and 73.3% A/A, in excellent agreement with those predicted for Hardy-Weinberg equilibrium.

Using an allele-specific odds ratio of 1.5 or greater as a practical level of significance (see Austin H et al., discussed above), the following observations can be made.

For breast cancer among black women, the odds ratio for the C allele as a risk factor was 3.9 (95% CI, 1.1-13). The odds ratio for the homozygote (CC) was a remarkable 14.7 (95% CI, 1.2-185). The heterozygote (CA genotype) had an odds ratio indistinguishable from 1 (odds ratio 0.5; 95% C.I. 0-5.3), suggesting that the C allele behaves as a recessive allele in this patient population.

For breast cancer among white women the odds ratio for the C allele as a risk factor was 2.3 (95% CI, 0.5-11). The odds ratio for the homozygote (CC genotype) was 4.8 (95% CI, 0.2-93). The heterozygote (CA genotype) had an odds ratio indistinguishable from 1 (odds ratio 1.2; 95% C.I. 0.1-14), suggesting that the C allele behaves as a recessive allele in this patient population.

For lung cancer in white men the odds ratio for the C allele as a risk factor was 5.1 (95% CI, 1.3-20). The C allele displayed a dosage effect, with the heterozygote (AC) having an odds ratio of 5.4 (95% CI, 0.7-42), and the homozygote (CC) an odds ratio of 7.2 (95% CI, 0.5-97). These data are consistent with a dominant action of the C allele, since one copy is sufficient to increase the odds ratio from 1 (for the AA homozygote) to 5.4 (for the AC heterozygote). The increase to 7.2 represents less than an additive effect of the allele, since 5.4 + 5.4 -1= 9.8 >7.2. These data are consistent with the C allele behaving as a dominant allele with interaction on less than an additive model.

For lung cancer in white women the odds ratio for the C allele (the novel SNP) as a risk factor was 3.5 (95% CI, 0.8-15). The CC homozygote surprisingly had a lower odds ratio, 0, than the heterozygote [9.5; 95% C.I., 1.3-71.0], suggesting that the C allele behaves other than as a simple recessive or dominant allele. The C allele may be

5   codominant with the A allele.

For prostate cancer among black men the odds ratio for the reference A allele as a risk factor was 2.3 (95% CI, 0.6-9.3). The odds ratio for the homozygote (AA genotype) was 3.7[H] (95% CI, 0.4-34). The heterozygote (AC genotype) had an odds ratio of 2.3[H] (95% CI, 0.2-22). The A allele therefore behaves as a dominant allele, with an additive

10   effect of increased allele dosage. The effect of the A allele on disease is as expected for an additive model (3.7 ~ 2.3 + 2.3 −1).

For prostate cancer in white men the odds ratio for the C allele (the novel SNP) as a risk factor was 6.0 (95% CI, 1.5-25). The CC homozygote surprisingly had a lower odds ratio (6.0; 95% CI, 0.3-124) than the heterozygote (15.0; 95% C.I. 1.9-116), suggesting

15   that the C allele behaves other than as a simple dominant or recessive allele. The C allele may be codominant with the A allele.

For hypertension among white men the odds ratio for the reference A allele as a risk factor was 2.2 (95% CI, 0.2-21). The odds ratio for the homozygote (AA genotype) was virtually the same (1.7[H]; 95% CI, 0.2-17) as that for the AC heterozygote (1.8[H]; 95%

20   CI, 0.1-26). These data indicate that the A allele behaves as a simple dominant allele, with no additional effect of a second copy of the allele.

For ESRD due to hypertension among black men the odds ratio for the reference A allele as a risk factor was 2.5 (95% CI, 0.4-15.2). The odds ratio for the homozygote (AA genotype) was 4.1[H] (95% CI, 0.4-42), and that for the AC heterozygote was essentially the

25   same [5.0[H] (95% CI, 0.4-60)]. These data suggest that the A allele behaves as a dominant allele.

For ESRD due to hypertension among black women the odds ratio for the C allele as a risk factor was 2.9 (95% CI, 0.5-17.2). The heterozygote (AC) had an odds ratio of 0, whereas the CC homozygote displayed an odds ratio of 7.9[H] (95% CI, 0.8-82). These data

30   are consistent with a recessive action of the C allele.

For ESRD due to hypertension among white men the odds ratio for the A allele as a risk factor was 3.5 (95% CI, 0.3-42.8). The odds ratio for the AC heterozygous genotype was 4.0 (95% CI, 0.3-55.5), and for the AA homozygous genotype was 1.9[H] (95% CI, 0.1-34). The A allele appears to be codominant with the C allele.

For ESRD due to hypertension among white women the odds ratio for the A allele was 2.0^H (95% CI, 0.2-18.8), relative to white women with hypertension but no renal disease. The odds ratios for both the AC heterozygote and the AA homozygote were only 1.0 after the Haldane's correction, shedding no light on the mechanism of action of the A

5   allele.

For NIDDM among black men, the odds ratio for the A allele at this locus was 2.3 (95% CI, 0.6-9.3). The odds ratio for the heterozygote was 2.3^H (95% CI, 0.2-22), and for the AA homozygote was 3.7^H (95% CI, 0.4-34). These data suggest that the A allele behaves as a dominant allele, with an additive effect of allele dosage (2.3 + 2.3 - 1 ~ 3.7).

10   For NIDDM among white men, the odds ratio for the C allele at this locus was 3.6 (95% CI, 0.9-14.4). The odds ratio for the heterozygote was 2.6 (95% CI, 0.3-22), and for the CC homozygote was 5.1 (95% CI, 0.4-66). These data suggest that the C allele behaves as a dominant allele, with more than an additive effect of allele dosage (2.6 + 2.6 - 1 < 5.1).

15   For NIDDM among white women, the odds ratio for the C allele at this locus was 2.3 (95% CI, 0.6-8.8). The odds ratio for the heterozygote was 2.4 (95% CI, 0.4-14), and for the CC homozygote was 3.2 (95% CI, 0.2-59). These data suggest that the C allele behaves as a dominant allele, with approximately an additive effect of allele dosage (2.4 + 2.4 - 1 = 3.8 ~ 3.2).

20   For ESRD due to NIDDM among black women, the odds ratio for the A allele at this locus was 1.6 (95% CI, 0.3-7.4). The odds ratio for the heterozygote was 3.9^H (95% CI, 0.3-46), and for the AA homozygote was 3.7^H (95% CI, 0.3-42). These data suggest that the A allele behaves as a classic dominant allele, with no effect of allele dosage.

For ESRD due to NIDDM among white men, the odds ratio for the A allele at this

25   locus was 2.6 (95% CI, 0.5-15.2). The odds ratio for the heterozygote was 5.0^H (95% CI, 0.4-60), and for the AA homozygote was 4.3^H (95% CI, 0.4-42). These data suggest that the A allele behaves as a classic dominant allele, with no effect of allele dosage.

For ESRD due to NIDDM among white women, the odds ratio for the A allele at this locus was 7.5 (95% CI, 0.9-62.1). The odds ratio for the heterozygote was

30   indistinguishable from 1, and for the AA homozygote was 2.5^H (95% CI, 0.2-28). These data suggest that the A allele behaves as a recessive allele.

For Caucasians with a history of alcohol abuse the odds ratio for the C allele was 1.7 (95% CI, 0.7 - 3.9). The odds ratio for the homozygote (C/C) was 1.6 (95% CI, 0.2 - 10.5), while the odds ratio for the heterozygote (C/ A) was 2.1 (95% CI, 0.7 - 6.5). These

data suggest that the C allele acts in a co-dominant manner in this patient population. These data further suggest that the EDN-1 gene is significantly associated with alcohol abuse in Caucasians, i.e. abnormal activity of the EDN-1 gene predisposes Caucasians to alcohol abuse.

5   For Caucasians with anxiety the odds ratio for the C allele was 1.6 (95% CI, 0.6 - 3.8). The odds ratio for the homozygote (C/C) was 0, while the odds ratio for the heterozygote (C/A) was 3.3 (95% CI, 1.1 - 10.3). These data suggest that the C allele acts in a co-dominant manner in this patient population. These data further suggest that the EDN-1 gene is significantly associated with anxiety in Caucasians, i.e. abnormal activity

10   of the EDN-1 gene predisposes Caucasians to anxiety.

For Caucasians with ASPVD due to NIDDM the odds ratio for the A allele was 1.8 (95% CI, 0.7- 4.8) , compared to Caucasians with NIDDM only. The odds ratio for the homozygote (A/A) was 0.2 (95% CI, 0 - 1.9), while the odds ratio for the heterozygote (C/A) was 2.0 (95% CI, 0.4 -9.4). These data suggest that the A allele acts in a co-

15   dominant manner in this patient population. These data further suggest that the EDN-1 gene is significantly associated with ASPVD due to NIDDM in Caucasians, i.e. abnormal activity of the EDN-1 gene predisposes Caucasians to ASPVD due to NIDDM.

For African-Americans with colon cancer the odds ratio for the A allele was 1.9 (95% CI, 0.8 - 4.7). Data were not sufficient to generate genotypic odds ratios of 1.5 or

20   greater. These data further suggest that the EDN-1 gene is significantly associated with colon cancer in African-Americans, i.e. abnormal activity of the EDN-1 gene predisposes African-Americans to colon cancer.

For African-Americans with COPD the odds ratio for the A allele was 5.3 (95% CI, 1.5 - 18.5). Data were not sufficient to generate genotypic odds ratios of 1.5 or

25   greater. These data further suggest that the EDN-1 gene is significantly associated with COPD in African-Americans, i.e. abnormal activity of the EDN-1 gene predisposes African-Americans to COPD .

For African-Americans with diabetic cardiomyopathy the odds ratio for the C allele was 2.9 (95% CI, 1 - 8.3) , compared to African-Americans with MI due to NIDDM.

30   The odds ratio for the homozygote (C/C) was 9.6[H] (95% CI, 0.2 - 574.5), while the odds ratio for the heterozygote (C/A) was 1.8 (95% CI, 0.5 - 6.6).These data suggest that the C allele acts in a dominant manner in this patient population. These data further suggest that the EDN-1 gene is significantly associated with diabetic cardiomyopathy in African-Americans, i.e. abnormal activity of the EDN-1 gene predisposes African-Americans to

diabetic cardiomyopathy .

For Caucasians with diabetic cardiomyopathy the odds ratio for the A allele was 1.6 (95% CI, 0.6- 4.7) , compared to Caucasians with MI due to NIDDM. Data were not sufficient to generate genotypic odds ratios of 1.5 or greater. These data further suggest that the EDN-1 gene is significantly associated with diabetic cardiomyopathy in Caucasians, i.e. abnormal activity of the EDN-1 gene predisposes Caucasians to diabetic cardiomyopathy.

For Caucasians with ESRD due to IDDM the odds ratio for the A allele was 2.7 (95% CI, 0.7 - 10). Data were not sufficient to generate genotypic odds ratios of 1.5 or greater. These data further suggest that the EDN-1 gene is significantly associated with ESRD due to IDDM in Caucasians, i.e. abnormal activity of the EDN-1 gene predisposes Caucasians to ESRD due to IDDM .

For African-Americans with ESRD due to IDDM the odds ratio for the C allele was 1.6 (95% CI, 0.6 - 3.9), compared to African-Americans with NIDDM only. The odds ratio for the homozygote (C/C) was 2.0 (95% CI, 0.3 - 14), while the odds ratio for the heterozygote (C/A) was 1.7 (95% CI, 0.5 - 6.1). These data suggest that the C allele acts in a dominant manner in this patient population with a less than additive effect of allele dosage [2 < 3.4 = (1.7 + 1.7 - 1.0)]. (Goldstein et al., *Monogr. Natl. Cancer Inst*; 26:49-54, 1999). These data further suggest that the EDN-1 gene is significantly associated with ESRD due to IDDM in African-Americans, i.e. abnormal activity of the EDN-1 gene predisposes African-Americans to ESRD due to IDDM .

For African-Americans with hypertensive cardiomyopathy the odds ratio for the A allele was 2.1 (95% CI, 0.7 – 6.0), compared to African-Americans with MI due to HTN. Data were not sufficient to generate genotypic odds ratios of 1.5 or greater. These data further suggest that the EDN-1 gene is significantly associated with hypertensive cardiomyopathy in African-Americans, i.e. abnormal activity of the EDN-1 gene predisposes African-Americans to hypertensive cardiomyopathy .

For Caucasians with hypertensive cardiomyopathy the odds ratio for the A allele was 2.5 (95% CI, 0.8 - 7.8) , compared to Caucasians with MI due to HTN. Data were not sufficient to generate genotypic odds ratios of 1.5 or greater. These data further suggest that the EDN-1 gene is significantly associated with hypertensive cardiomyopathy in Caucasians, i.e. abnormal activity of the EDN-1 gene predisposes Caucasians to hypertensive cardiomyopathy .

For Caucasians with CVA due to NIDDM the odds ratio for the A allele was 3.0 (95% CI, 1 - 9.4) , compared to Caucasians with NIDDM only . The odds ratio for the homozygote (A/A) was 0 , while the odds ratio for the heterozygote (C/A) was 1.5 (95% CI, 0.3 - 7.2).These data suggest that the A allele acts in a manner in this patient

5   population . These data further suggest that the EDN-1 gene is significantly associated with CVA due to NIDDM in Caucasians, i.e. abnormal activity of the EDN-1 gene predisposes Caucasians to CVA due to NIDDM .

For African-Americans with MI due to NIDDM the odds ratio for the A allele was 2.3 (95% CI, 0.8 - 7), compared to African-Americans with NIDDM only . Data were not

10   sufficient to generate genotypic odds ratios of 1.5 or greater. These data further suggest that the EDN-1 gene is significantly associated with MI due to NIDDM in African-Americans, i.e. abnormal activity of the EDN-1 gene predisposes African-Americans to MI due to NIDDM .

For African-Americans with seizure disorder the odds ratio for the C allele was 1.8

15   (95% CI, 0.9 - 3.9). The odds ratio for the homozygote (C/C) was 3.0 (95% CI, 0.6 - 14.9), while the odds ratio for the heterozygote (C/A) was 1.9 (95% CI, 0.6 - 5.8).These data suggest that the C allele acts in a dominant manner in this patient population with a greater than additive effect of allele dosage [3 > 3.8 = (1.9 + 1.9 - 1.0)]. (Goldstein et al., *Monogr. Natl. Cancer Inst.*; 26:49-54, 1999). These data further suggest that the EDN-1

20   gene is significantly associated with seizure disorder in African-Americans, i.e. abnormal activity of the EDN-1 gene predisposes African-Americans to seizure disorder.

For Caucasians with seizure disorder the odds ratio for the A allele was 1.7 (95% CI, 0.6 - 5.1). Data were not sufficient to generate genotypic odds ratios of 1.5 or greater. These data further suggest that the EDN-1 gene is significantly associated with seizure

25   disorder in Caucasians, i.e. abnormal activity of the EDN-1 gene predisposes Caucasians to seizure disorder.

According to MatInspector (GENOMATIX; see above for URL and reference), the C2657→A SNP disrupts a binding site for CEBPB (CCAAT/enhancer binding protein-beta; Quandt K et al., *Nucleic Acids Res.*, 23: 4878-4884 (1995)). CEBPB binds to a core

30   sequence of four nucleotides, GMAA, in an overall sequence of 14 nucleotides (ref. Akira, S. et al. A nuclear factor for IL-6 expression (NF-IL6) is a member of a C/EBP family. EMBO J. 1990 Jun:9(6):1897-1906). CEBPB_01 binding sites, which occur on average 2.07 times per 1000 base pairs of random genomic sequence in vertebrates, are predicted to occur at positions 2647 to 2660 on the (+) strand of reference sequence J05008.1

(matrix score 0.952, with 1.0 being an identical match), as well as from position 2670 to 2657 on the (-) strand (matrix score 0.891 out of a possible 1.0). In neither case, however, does the C2657→A SNP alter a nucleotide critical for binding.

5

Table 17

Gene	Region	Location	Reference Type	Variant	SEQ ID
EDN-1	Promoter	2239	T	G	1
		2657	A	C	1

Conclusion

In light of the detailed description of the invention and the examples presented above, it can be appreciated that the several aspects of the invention are achieved.

It is to be understood that the present invention has been described in detail by way of illustration and example in order to acquaint others skilled in the art with the invention, its principles, and its practical application. Particular formulations and processes of the present invention are not limited to the descriptions of the specific embodiments presented, but rather the descriptions and examples should be viewed in terms of the claims that follow and their equivalents. While some of the examples and descriptions above include some conclusions about the way the invention may function, the inventor does not intend to be bound by those conclusions and functions, but puts them forth only as possible explanations.

It is to be further understood that the specific embodiments of the present invention as set forth are not intended as being exhaustive or limiting of the invention, and that many alternatives, modifications, and variations will be apparent to those of ordinary skill in the art in light of the foregoing examples and detailed description. Accordingly, this invention is intended to embrace all such alternatives, modifications, and variations that fall within the spirit and scope of the following claims.

5   What is claimed is:

1. A method for diagnosing a genetic susceptibility for a disease, condition, or disorder in a subject comprising:

    obtaining a biological sample containing nucleic acid from said subject; and

    analyzing said nucleic acid to detect the presence or absence of a single nucleotide

10   polymorphism in the EDN-1 gene, wherein said single nucleotide polymorphism is associated with a genetic predisposition for a disease, condition or disorder selected from the group consisting of hypertension, end stage renal disease due to hypertension, non-insulin dependent diabetes mellitus, end stage renal disease due to non-insulin dependent diabetes mellitus, lung cancer, breast cancer, prostate cancer, colon cancer,

15   atherosclerotic peripheral vascular disease due to hypertension, cerebrovascular accident due to hypertension, cataracts due to hypertension, cardiomyopathy with hypertension, myocardial infarction due to hypertension, atherosclerotic peripheral vascular disease due to non-insulin dependent diabetes mellitus, cerebrovascular accident due to non-insulin dependent diabetes mellitus, ischemic cardiomyopathy,

20   ischemic cardiomyopathy with non-insulin dependent diabetes mellitus, myocardial infarction due to non-insulin dependent diabetes mellitus, atrial fibrillation without valvular disease, alcohol abuse, anxiety, asthma, chronic obstructive pulmonary disease, cholecystectomy, degenerative joint disease, end stage renal disease and frequent de-clots, end stage renal disease due to focal segmental glomerular sclerosis,

25   end stage renal disease due to insulin dependent diabetes mellitus, or seizure disorder.

2. The method of claim 1, wherein the gene EDN-1 comprises SEQ ID NO: 1.

5

3. The method of claim 1, wherein said nucleic acid is DNA, RNA, cDNA or mRNA.

5

4. The method of claim 2, wherein said single nucleotide polymorphism is located at position 2239 or 2657 of SEQ ID NO: 1.

5. The method of claim 4, wherein said single nucleotide polymorphism is selected from the group consisting of T2239->G and C2657->A and the complements thereof namely A2239->C and G2657->T.

6. The method of claim 1, wherein said analysis is accomplished by sequencing, mini sequencing, hybridization, restriction fragment analysis, oligonucleotide ligation assay or allele specific PCR.

7. An isolated polynucleotide comprising at least 10 contiguous nucleotides of SEQ ID NO: 1, or the complement thereof, and containing at least one single nucleotide polymorphism at position 2239 or 2657 of SEQ ID NO: 1 wherein said at least one single nucleotide polymorphism is associated with a disease, condition or disorder

5   selected from the group consisting of hypertension, end stage renal disease due to hypertension, non-insulin dependent diabetes mellitus, end stage renal disease due to non-insulin dependent diabetes mellitus, lung cancer, breast cancer, prostate cancer, colon cancer, atherosclerotic peripheral vascular disease due to hypertension, cerebrovascular accident due to hypertension, cataracts due to hypertension,

10   cardiomyopathy with hypertension, myocardial infarction due to hypertension, atherosclerotic peripheral vascular disease due to non-insulin dependent diabetes mellitus, cerebrovascular accident due to non-insulin dependent diabetes mellitus, ischemic cardiomyopathy, ischemic cardiomyopathy with non-insulin dependent diabetes mellitus, myocardial infarction due to non-insulin dependent diabetes

15   mellitus, atrial fibrillation without valvular disease, alcohol abuse, anxiety, asthma, chronic obstructive pulmonary disease, cholecystectomy, degenerative joint disease, end stage renal disease and frequent de-clots, end stage renal disease due to focal segmental glomerular sclerosis, end stage renal disease due to insulin dependent diabetes mellitus, or seizure disorder.

8. The isolated polynucleotide of claim 7, wherein at least one single nucleotide polymorphism is selected from the group consisting of T2239->G and C2657->A and the complements thereof namely A2239->C and G2657->T.

9. The isolated polynucleotide of claim 7, wherein said at least one single nucleotide polymorphism is located at the 3' end of said nucleic acid sequence.

10. The isolated polynucleotide of claim 7, further comprising a detectable label.

11. The isolated nucleic acid sequence of claim 10, wherein said detectable label is selected from the group consisting of radionuclides, fluorophores or fluorochromes, peptides, enzymes, antigens, antibodies, vitamins or steroids.

12. A kit comprising at least one isolated polynucleotide of at least 10 contiguous nucleotides of SEQ ID NO: 1 or the complement thereof, and containing at least one single nucleotide polymorphism associated with a disease, condition, or disorder
5   selected from the group consisting of hypertension, end stage renal disease due to hypertension, non-insulin dependent diabetes mellitus, end stage renal disease due to non-insulin dependent diabetes mellitus, lung cancer, breast cancer, prostate cancer, colon cancer, atherosclerotic peripheral vascular disease due to hypertension, cerebrovascular accident due to hypertension, cataracts due to hypertension,
10   cardiomyopathy with hypertension, myocardial infarction due to hypertension, atherosclerotic peripheral vascular disease due to non-insulin dependent diabetes mellitus, cerebrovascular accident due to non-insulin dependent diabetes mellitus, ischemic cardiomyopathy, ischemic cardiomyopathy with non-insulin dependent diabetes mellitus, myocardial infarction due to non-insulin dependent diabetes
15   mellitus, atrial fibrillation without valvular disease, alcohol abuse, anxiety, asthma, chronic obstructive pulmonary disease, cholecystectomy, degenerative joint disease, end stage renal disease and frequent de-clots, end stage renal disease due to focal segmental glomerular sclerosis, end stage renal disease due to insulin dependent diabetes mellitus, and seizure disorder; and instructions for using said polynucleotide
20   for detecting the presence or absence of said at least one single nucleotide polymorphism in said nucleic acid.

13. The kit of claim 12 wherein said at least one single nucleotide polymorphism is located at position 2239 or 2657 of SEQ ID NO: 1.

14. The kit of claim 13 wherein said at least one single nucleotide polymorphism selected from the group consisting of T2239->G and C2657->A and the complements thereof namely A2239->C and G2657->T.

15. The kit of claim 12, wherein said single nucleotide polymorphism is located at the 3' end of said polynucleotide.

16. The kit of claim 12, wherein said polynucleotide further comprises at least one detectable label.

17. The kit of claim 16, wherein said label is chosen from the group consisting of radionuclides, fluorophores or fluorochromes, peptides enzymes, antigens, antibodies, vitamins or steroids.

18. A kit comprising at least one polynucleotide of at least 10 contiguous nucleotides of SEQ ID NO: 1 or the complement thereof, wherein the 3' end of said polynucleotide is immediately 5' to a single nucleotide polymorphism site associated with a genetic predisposition to disease, condition, or disorder selected from the group consisting of hypertension, end stage renal disease due to hypertension, non-insulin dependent diabetes mellitus, end stage renal disease due to non-insulin dependent diabetes mellitus, lung cancer, breast cancer, prostate cancer, colon cancer, atherosclerotic peripheral vascular disease due to hypertension, cerebrovascular accident due to hypertension, cataracts due to hypertension, cardiomyopathy with hypertension, myocardial infarction due to hypertension, atherosclerotic peripheral vascular disease due to non-insulin dependent diabetes mellitus, cerebrovascular accident due to non-insulin dependent diabetes mellitus, ischemic cardiomyopathy, ischemic cardiomyopathy with non-insulin dependent diabetes mellitus, myocardial infarction due to non-insulin dependent diabetes mellitus, atrial fibrillation without valvular disease, alcohol abuse, anxiety, asthma, chronic obstructive pulmonary disease, cholecystectomy, degenerative joint disease, end stage renal disease and frequent de-clots, end stage renal disease due to focal segmental glomerular sclerosis, end stage renal disease due to insulin dependent diabetes mellitus, and seizure disorder; and instructions for using said polynucleotide for detecting the presence or absence of said single nucleotide polymorphism in a biological sample containing nucleic acid.

19. The kit of claim 18, wherein said single nucleotide polymorphism site is located at position 2239 or 2657 of SEQ ID NO: 1.

20. The kit of claim 19, wherein said at least one polynucleotide further comprises a detectable label.

21. The kit of claim 20, wherein said detectable label is chosen from the group consisting of radionuclides, fluorophores or fluorochromes, peptides, enzymes, antigens, antibodies, vitamins or steroids.

22. A method for treatment or prophylaxis in a subject comprising:

obtaining a sample of biological material containing nucleic acid from a subject;

analyzing said nucleic acid to detect the presence or absence of at least one single
5   nucleotide polymorphism in SEQ ID NO: 1 or the complement thereof associated with a disease, condition, or disorder selected from the group consisting of hypertension, end stage renal disease due to hypertension, non-insulin dependent diabetes mellitus, end stage renal disease due to non-insulin dependent diabetes mellitus, lung cancer, breast cancer, prostate cancer, colon cancer, atherosclerotic peripheral vascular disease
10   due to hypertension, cerebrovascular accident due to hypertension, cataracts due to hypertension, cardiomyopathy with hypertension, myocardial infarction due to hypertension, atherosclerotic peripheral vascular disease due to non-insulin dependent diabetes mellitus, cerebrovascular accident due to non-insulin dependent diabetes mellitus, ischemic cardiomyopathy, ischemic cardiomyopathy with non-insulin
15   dependent diabetes mellitus, myocardial infarction due to non-insulin dependent diabetes mellitus, atrial fibrillation without valvular disease, alcohol abuse, anxiety, asthma, chronic obstructive pulmonary disease, cholecystectomy, degenerative joint disease, end stage renal disease and frequent de-clots, end stage renal disease due to focal segmental glomerular sclerosis, end stage renal disease due to insulin dependent
20   diabetes mellitus, and seizure disorder; and treating said subject for said disease, condition or disorder.

23. The method of claim 22 wherein said nucleic acid is selected from the group consisting of DNA, cDNA, RNA and mRNA.

24. The method of claim 22, wherein said at least one single nucleotide polymorphism is located at position 2239 or 2657 of SEQ ID NO: 1.

25. The method of claim 22 wherein said at least one single nucleotide polymorphism is selected from the group consisting of T2239->G and C2657->A and the complements thereof namely A2239->C and G2657->T.

26. The method of claim 22 wherein said treatment counteracts the effect of said at least one single nucleotide polymorphism detected.

Figure 1

```
gatatcctat taatacagag atacagaaag aaatacataa aaaatagttt tatcaaatac   60
tttccagcat tcaagtgtag cctcaaaagc aagaataggc caggagtggt ggctcacgct  120
gtaatccaca gcactgtggg aggccaaggt aagaggattg cttgaggcca ggatttcaag  180
accagcctag gcaacatagt gagatcccta tctctacgaa aaaattttaa aacttagctg  240
ggcatggtgc ttgagcctgt tgtcccagct actcaggagg tgaagtagga gtgtcacttg  300
agcccaggag gttgaggctg cagtgagcta taactgcacc actgcactcc agccttggag  360
acagagtgag accctgtccc caaaaaaatt aaaattgaga aaaaaaaaaa ggcaagaaca  420
gccacagcaa actttctatt ggggaaaaaa aaaaatcctc ctctttacat ctctcccttc  480
cttcccttcc ctttctgaga gtgactgtgg ccaaaaggag catttctccc ctgcagtcct  540
ctgaggggtg gggtggggct atgaagctat ccttcatatt cactcctttg tccagctctt  600
ttcacctcta gttcttctcc ccgcatctct gtctagcagt gccttaagtg gaggaggggt  660
gggggcatca agcttgtaaa actggtttgt tggggttctc cttctcccct catttcttga  720
ttcttgggaa aatgtcttgc tgggaggctg ccctagctg ccttctgtgg  780
```

(continues on 2/5)

```
gcttgaatgg ggcttccctc tgcccctaca ggaggaaaag ggagctgctg ccagagggag    840
aaatggagag atggacagag aaggcaggtg ccacccctcg cccctgacac acaaagaaaa    900
agacacggaa attctctctc tctcttctct tctcctatct ctctctctct ccctctctct    960
ctctctctct ctctctctca cacacacaca cacacacaca caggcgcgcg              1020
ccgcgcgcgc aggcacacgt cttgcaaatt caggattcaa agagacaggg gcaccattat   1080
atttggcacg gtggggcctt ccaggtctga aatcctgcat tcttccttac tatttacttt   1140
ccccgagctc gagaagggcc aggtgtgggc ggatggctgg ccacgttttg tgtttccaat   1200
tcatattcac gggatgacac agacggggcg tggtgagtgc tgttggaggc gcttgggcag   1260
tttcattttg ccccacttct ccacctgaag gctgggcgtt gctggaacct gcaggggcag   1320
cctcagcaag gtggggtggc gtggagtggg gtgggagaag ggactccagc tgaagtagaa   1380
cccaggctgg acctgagaat attgggggag gcatgggcgg tggtttccgg gtaggggcct   1440
tgaggacatg ttggtcctga ctgttgtcag tgtttggtca aagttgccaa aaggttaaaa   1500
aaaaaaaagt aggggggagtc cctgccaaga catatttccc aggccacctt tcttccgcgg  1560
gagtgttggg ggggaggcgc tgcttggaac ctgtgaatgt gacatcagct ctcctctcct   1620
ctcccaaggt cggctttgga gagggaggtc aggcaccct tgcctggcac aggcacgctg    1680
```

(continues on 3/5)

```
gcttccggct cagtgccgcc tgctctccgg gagctgtgcg ctccctgggc cccggggcta   1740
ggctgaggta agcgcacagc ggaggccagg cgcgccggca gaggcctggg ggatagggtg   1800
gaggcatctc tgggtgtggg tgtgggtgtg ggtgtgggag ggagagttct tgcctctctc   1860
tctccatct  ccaactcttg cttcagtggc tcttttagag gatgcatgtc attatggacc   1920
tgtcgctgcc actgtccctg ttcccccagc tgtgacttcg agggaggtct ggggatctga   1980
gtctgtccaa acccacggct ttgctgttgg gataaaaact gtcctttga  ttttagaagg   2040
aggaggaaa  aaaggtttcc cagcatgtgt gttgtgccag tcttggaaat tcatccgtgc   2100
ttgaattcca ccctccatcc ccagaaaaac tggagtaaaa caaaaagagg agatggacaa   2160
agtgtgtatt tgatggcatc ccctgggaag agactctaaa tttatcccat aggtcttact   2220
gggccactgt gagcgctttg gtggagaaca aacaaaaatt ctgggtgctc agttgtctaa   2280
cctgaaaaat gggactagcg gaaaaagcca atgtgttcca tgcacctttt gctttcttta   2340
ttaaggcatg atgtcacctg tacagtaact gccctgtgtg tacttcaggg ggggatttca   2400
aggttagata gacaggaaat tgtttgaaa  atgtaaacac attattaaat gtgaagtatt   2460
atctgattcc ttgttcgaat ggcatttcct tctcagcacc accttccttg catattcact   2520
taaccttgta caagaacacc tttttgccct aaatgaagac accccccaa  aaaaaagagt   2580
```

(continues on 4/5)

```
cccagaaaat atgtccctgc ttgtgcggga ataaatagaa tattctgagg tgcattcctc    2640

cttcctatgt taggcaacat tccttgaccc tcctcggccc ccaagccagg ttgcgttttt    2700

ttctgccatt tagaagggtt ttcctttttg tcctagtaaa acatcagccc ctgtagctct    2760

tcatctcccc ctggtgttct tctcccgcca tgtcttaaga ttggtggcac cgaccaatct    2820

taagatttaa gttctgtgtg aaaaacacct ttgcttttca atcagtttat cagcctcctc    2880

cgcaggggaa tgtggacaca caaaagaact tatcggggct tctcatcagt gataggggaaa    2940

agactggcat gtgcctaaac gagctctgat gttatttta agctccctt cttgccaatc    3000

cctcacggat ctttctccga tagatgcaaa gaacttcagc aaaaaagacc cgcaggaagg    3060

ggcttgaaga gaaaagtacg ttgatctgcc aaaatagtct gacccccagt agtgggcagt    3120

gacgagggag agcattccct tgtttgactg agactagaat cggagagaca taaaaggaaa    3180

atgaagcgag caacaattaa aaaaaattcc ccgcacacaa caatacaatc tatttaaact    3240

gtggctcata ctttccatac caatggtatg acttttttttc tggagtcccc tcttctgatt    3300

cttgaactcc ggggctggca gcttgcaaag gggaagcgga ctccagcact gcacgggcag    3360

gtttagcaaa ggtctctaat gggtattttc tttttcttag ccctgccccc gaattgtcag    3420

acggcgggcg tctgcttctg aagttagcag tgatttcctt tcgggcctgg cttatctccg    3480

(continues on 5/5)
```

gctgcacgtt gcctgttggt gactaataac acaataacat tgtctggggc tggaataaag   3540

tcggagctgt ttaccccac tctaataggg gttcaatata aaaagccggc agagagctgt   3600

ccaagtc aga cgc gcc tct gca tct gcg cca ggc gaa cgg gtc ctg cgc   3649

1

SEQUENCE LISTING

<110> DzGenes LLC

<120> ENDOTHELIN-1 PROMOTER POLYMORPHISM

<130> DZG 2198

<150> US 60/231,672
<151> 2000-09-11

<150> US 60/243,814
<151> 2000-10-27

<160> 5

<170> PatentIn version 3.0

<210> 1
<211> 12459
<212> DNA
<213> Homo sapiens

<220>
<221> repeat_region
<222> (98)..(383)

<220>
<221> protein_bind
<222> (739)..(745)
<223> acute phase reactant regulatory element

<220>
<221> misc_feature
<222> (979)..(1039)
<223> Z-DNA region; putative

<220>
<221> protein_bind
<222> (2183)..(2188)
<223> acute phase reactant regulatory element

<220>
<221> protein_bind
<222> (2951)..(2958)
<223> TPA/JUN

<220>
<221> CAAT_signal
<222> (3510)..(3515)

<220>
<221> exon

```
<222>    (3608)..(3939)

<220>
<221>  sig_peptide
<222>    (3876)..(3938)

<220>
<221>  Intron
<222>    (3940)..(5584)

<220>
<221>  protein_bind
<222>    (4612)..(4618)

<220>
<221>  exon
<222>    (5585)..(5753)

<220>
<221>  Intron
<222>    (5754)..(7182)

<220>
<221>  exon
<222>    (7183)..(7338)

<220>
<221>  Intron
<222>    (7339)..(7502)

<220>
<221>  exon
<222>    (7503)..(7646)

<220>
<221>  Intron
<222>    (7647)..(9211)

<220>
<221>  protein_bind
<222>    (8205)..(8218)

<220>
<221>  protein_bind
<222>    (8260)..(8266)

<220>
<221>  exon
<222>    (9212)..(10443)

<220>
<221>  protein_bind
<222>    (9392)..(9398)

<220>
<221>  protein_bind
<222>    (10751)..(10764)
```

```
<220>
<221>  protein_bind
<222>  (3241)..(3248)
<223>  TPA/JUN


<220>
<221>  protein_bind
<222>  (3316)..(3328)
<223>  TPA/JUN


<220>
<221>  protein_bind
<222>  (3499)..(3505)
<223>  TPA/JUN


<220>
<221>  TATA_signal
<222>  (3577)..(3582)

<400>  1
gatatcctat taatacagag atacagaaag aaatacataa aaaatagttt tatcaaatac       60

tttccagcat tcaagtgtag cctcaaaagc aagaataggc caggagtggt ggctcacgct      120

gtaatccaca gcactgtggg aggccaaggt aagaggattg cttgaggcca ggatttcaag      180

accagcctag gcaacatagt gagatcccta tctctacgaa aaaattttaa aacttagctg      240

ggcatggtgc ttgagcctgt tgtcccagct actcaggagg tgaagtagga gtgtcacttg      300

agcccaggag gttgaggctg cagtgagcta taactgcacc actgcactcc agccttggag      360

acagagtgag accctgtccc caaaaaaatt aaaattgaga aaaaaaaaa ggcaagaaca      420

gccacagcaa actttctatt ggggaaaaaa aaaaatcctc ctctttacat ctctcccttc      480

cttcccttcc ctttctgaga gtgactgtgg ccaaaaggag cattttcccc ctgcagtcct      540

ctgaggggtg gggtggggct atgaagctat ccttcatatt cactcctttg tccagctctt      600

ttcacctcta gttcttctcc ccgcatctct gtctagcagt gccttaagtg gaggaggggt      660

gggggcatca agcttgtaaa actggtttgt tggggttctc cttctcccct catttcttga      720

ttcttgggaa aatgtcttgc tgggaggctg cctggcgagt gccctagctg ccttctgtgg      780

gcttgaatgg ggcttccctc tgcccctaca ggaggaaaag ggagctgctg ccagagggag      840

aaatggagag atggacagag aaggcaggtg ccacccctcg cccctgacac acaaagaaaa      900

agacacggaa attctctctc tctcttctct tctcctatct ctctctctct ccctctctct      960

ctctctctct ctctctctca cacacacaca cacacacaca cacacacaca caggcgcgcg     1020
```

```
ccgcgcgcgc aggcacacgt cttgcaaatt caggattcaa agagacaggg gcaccattat    1080

atttggcacg gtggggcctt ccaggtctga aatcctgcat tcttccttac tatttacttt    1140

ccccgagctc gagaagggcc aggtgtgggc ggatggctgg ccacgttttg tgtttccaat    1200

tcatattcac gggatgacac agacggggcg tggtgagtgc tgttggaggc gcttgggcag    1260

tttcattttg ccccacttct ccacctgaag gctgggcgtt gctggaacct gcaggggcag    1320

cctcagcaag gtggggtggc gtggagtggg gtgggagaag ggactccagc tgaagtagaa    1380

cccaggctgg acctgagaat attggggagg gcatgggcgg tggtttccgg gtaggggcct    1440

tgaggacatg ttggtcctga ctgttgtcag tgtttggtca aagttgccaa aaggttaaaa    1500

aaaaaaaagt aggggagtc cctgccaaga catatttccc aggccacctt tcttccgcgg    1560

gagtgttggg ggggaggcgc tgcttggaac ctgtgaatgt gacatcagct ctcctctcct    1620

ctcccaaggt cggctttgga gagggaggtc agggcaccct tgcctggcac aggcacgctg    1680

gcttccggct cagtgccgcc tgctctccgg gagctgtgcg ctccctgggc cccggggcta    1740

ggctgaggta agcgcacagc ggaggccagg cgcgccggca gaggcctggg ggatagggtg    1800

gaggcatctc tgggtgtggg tgtgggtgtg ggtgtgggag ggagagttct tgcctctctc    1860

tctcccatct ccaactcttg cttcagtggc tcttttagag gatgcatgtc attatggacc    1920

tgtcgctgcc actgtccctg ttcccccagc tgtgacttcg agggaggtct ggggatctga    1980

gtctgtccaa acccacggct ttgctgttgg gataaaaact gtccttttga ttttagaagg    2040

aggagggaaa aaaggtttcc cagcatgtgt gttgtgccag tcttggaaat tcatccgtgc    2100

ttgaattcca ccctccatcc ccagaaaaac tggagtaaaa caaaaagagg agatggacaa    2160

agtgtgtatt tgatggcatc ccctgggaag agactctaaa tttatcccat aggtcttact    2220

gggccactgt gagcgctttg gtggagaaca aacaaaaatt ctgggtgctc agttgtctaa    2280

cctgaaaaat gggactagcg gaaaaagcca atgtgttcca tgcacctttt gctttctta    2340

ttaaggcatg atgtcacctg tacagtaact gccctgtgtg tacttcaggg ggggatttca    2400

aggttagata gacaggaaat tgttttgaaa atgtaaacac attattaaat gtgaagtatt    2460

atctgattcc ttgttcgaat ggcatttcct tctcagcacc accttccttg catattcact    2520

taaccttgta caagaacacc tttttgccct aaatgaagac acccccccaa aaaaaagagt    2580

cccagaaaat atgtccctgc ttgtgcggga ataaatagaa tattctgagg tgcattcctc    2640

cttcctatgt taggcaacat tccttgaccc tcctcggccc ccaagccagg ttgcgttttt    2700
```

ttctgccatt tagaagggtt ttcctttttg tcctagtaaa acatcagccc ctgtagctct   2760

tcatctcccc ctggtgttct tctcccgcca tgtcttaaga ttggtggcac cgaccaatct   2820

taagatttaa gttctgtgtg aaaaacacct ttgcttttca atcagtttat cagcctcctc   2880

cgcaggggaa tgtggacaca caaaagaact tatcggggct tctcatcagt gatagggaaa   2940

agactggcat gtgcctaaac gagctctgat gttatttta agctcccttt cttgccaatc   3000

cctcacggat ctttctccga tagatgcaaa gaacttcagc aaaaaagacc cgcaggaagg   3060

ggcttgaaga gaaaagtacg ttgatctgcc aaaatagtct gaccccagt agtgggcagt   3120

gacgagggag agcattccct tgtttgactg agactagaat cggagagaca taaaaggaaa   3180

atgaagcgag caacaattaa aaaaaattcc ccgcacacaa caatacaatc tatttaaact   3240

gtggctcata cttttcatac caatggtatg actttttttc tggagtcccc tcttctgatt   3300

cttgaactcc ggggctggca gcttgcaaag gggaagcgga ctccagcact gcacgggcag   3360

gtttagcaaa ggtctctaat gggtatttc tttttcttag ccctgccccc gaattgtcag   3420

acggcgggcg tctgcttctg aagttagcag tgatttcctt tcgggcctgg cttatctccg   3480

gctgcacgtt gcctgttggt gactaataac acaataacat tgtctggggc tggaataaag   3540

tcggagctgt ttaccccac tctaataggg gttcaatata aaaagccggc agagagctgt   3600

```
ccaagtc aga cgc gcc tct gca tct gcg cca ggc gaa cgg gtc ctg cgc    3649
        Arg Arg Ala Ser Ala Ser Ala Pro Gly Glu Arg Val Leu Arg
        1             5               10

ctc ctg cag tcc cag ctc tcc acc gcc gcg tgc gcc tgc aga cgc tcc    3697
Leu Leu Gln Ser Gln Leu Ser Thr Ala Ala Cys Ala Cys Arg Arg Ser
15              20              25              30

gct cgc tgc ctt ctc tcc tgg cag gcg ctg cct ttt ctc ccc gtt aaa    3745
Ala Arg Cys Leu Leu Ser Trp Gln Ala Leu Pro Phe Leu Pro Val Lys
            35              40              45

ggg cac ttg ggc tga agg atc gct ttg aga tct gag gaa ccc gca gcg    3793
Gly His Leu Gly     Arg Ile Ala Leu Arg Ser Glu Glu Pro Ala Ala
            50              55              60

ctt tga ggg acc tga agc tgt ttt tct tcg ttt tcc ttt ggg ttc agt    3841
Leu     Gly Thr     Ser Cys Phe Ser Ser Phe Ser Phe Gly Phe Ser
                    65              70              75

ttg aac ggg agg ttt ttg atc cct ttt ttt cag aat gga tta ttt gct    3889
Leu Asn Gly Arg Phe Leu Ile Pro Phe Phe Gln Asn Gly Leu Phe Ala
            80              85              90

cat gat ttt ctc tct gct gtt tgt ggc ttg cca agg agc tcc aga aac    3937
His Asp Phe Leu Ser Ala Val Cys Gly Leu Pro Arg Ser Ser Arg Asn
            95              100             105
```

ag gtaggcacgc tcgttgactt gtaagtctcg gaattacaag ttagtgtgtt   3989

cttatccacc ttcatgcttt tcttgcttct attttttcccc gttctttta tgactgcagc   4049

ttagagagca agtgtctgag aattattgct gaaacgtact ttaagtcttc tagtgtaaaa   4109

tgtaaaattc ctctactgaa tacaattagg tgcaattgac tataacatga cattaaaata   4169

acttatcgtt ttattattat tattccatta tgtgtttcct tggctttttaa aaaatgagaa 4229

gagtatggac atatacaatt tagtcaaatg tatgtttgta atatatgtgt ttatacaggt   4289

acacaggcca tataggaact taaatcttat ttaaacacta ttttaatagt gtgttaacgt   4349

gtaaaatatt taagcattcc agcttgaagc caaggaattg tatccagtcg ttcaagcaat   4409

gtatgttcag taaaatcacc tgcagagcaa aagtctgttg actaactacc gcctcccccg   4469

ccccccacc accccccgca ggcggtttct gggtgaagca gatgttttct ttaaaatttg   4529

tcatcattga ctttaggttt cttttggcag gtttttggca cccaaaacag tgtgagctct   4589

cttttcagct ttattcacct gtgctgggag gggagctagg ataattcttg gctgccgaag   4649

gatttaggca gtgcgtgtgc atctgcccgg gtccccccg tttttagggt cagtgcactt   4709

tttttgtctt ttcgtgaccc tgactaaaga gaaaggatgt caagggaatg aaaatcctgg   4769

aatgtgtctg atcatttgaa atgtacaaaa ttgggcagat aagctgcatg gctaaattgt   4829

taggaggaag aggcaaggca gtagtggaga aggggaggc agtggatccc acacaagcct   4889

gatgcccagg gattcggaat tcaaaatccc cccagcctac cttcagtccc ctgacctgct   4949

tctcagcccc accttaggtc actggtttct atggagttac cctactgaat tgaatattga   5009

atagttaatt tctctctcca atcattttcc ccacctaatt ttgaaagata tacatcatct   5069

ggggtaccct gtgccctaca cagcatgtga agtggatggg tacccctaa agagagggtc   5129

atcctgaatg gggaagtggc cccaaagcta ggaataactg tgatttcttg tctttagtca   5189

tgtgccaatg ttaagtaagc ttcagtggat agtgctgtcc taccaagttc cttgtagaag   5249

ccagccggat tttcaacagg cagcattcca cagcatttcc ctgagcctgc ttcaagaggg   5309

gtgggggaag tcccttttca ggtgtttatc tcctctgcat ttgtgtaatc tccctgaagg   5369

tggataagcc aagggcatga gggggaggca aaaggtgaac tcatgttaag gagggaaaaa   5429

aataaagagc cctttttttct gtgtttcttg ctgatggcag gctgtgtgct tcatctgctt   5489

ttatctgctc tgctagctct gactctactg tgatccagca tgtctctcgg cgtttgagga   5549

gacatccccc actgacctgc tctttctctc cccag cag tct tag gcg ctg agc   5602
                                       Gln Ser   Ala Leu Ser

7

110

```
tca gcg cgg tgg gtg aga acg gcg ggg aga aac cca ctc cca gtc cac   5650
Ser Ala Arg Trp Val Arg Thr Ala Gly Arg Asn Pro Leu Pro Val His
    115             120                 125

cct ggc ggc tcc gcc ggt cca agc gct gct cct gct cgt ccc tga tgg   5698
Pro Gly Gly Ser Ala Gly Pro Ser Ala Ala Pro Ala Arg Pro     Trp
130             135                 140

ata aag agt gtg tct act tct gcc acc tgg aca tca ttt ggg tca aca   5746
Ile Lys Ser Val Ser Thr Ser Ala Thr Trp Thr Ser Phe Gly Ser Thr
145             150                 155                 160

ctc ccg a gtaagtctct agagggcatt gtaaccctat tcattcatta gcgctggctc   5803
Leu Pro
```

```
cactggagcc cagttttaga gtttcttttc tagggactct gaaggtagtc cttctaacac   5863

catccaagtg cctcagtggg gacagtttcc ctctattcct gaaaataacg acagcttcgt   5923

tcttagcaac caaggggagg gtcttctgag gccccgtagc tcaggctact catgatggga   5983

caagcaggag gccactgcac gtttcaaatg aggaactttc agtgagaggg cctcaggggg   6043

acactctcac agtggcatct gatggggttt cgggaataat tgccgaggtc agatgtgggt   6103

tagtgcaacc tgtgcttctc atgggagggt ggagactgag aggcagaagt gatgatatag   6163

agggttagaa tcacttaatt ttagttacag aaaaacctag gctcaaagtg ttgaagccat   6223

ttgtgcagga gtgagtttgt agcagagcta gaactggagc ccggatttcc tttgctgcta   6283

tattttccct ttagaaatgc ccatttcaga actgaaatag aaatactgtc cataggcttc   6343

tctttcacct acagagaaga aaagcagatt cctccttct gccctggaca ctagttcatc   6403

atctgtcgga agcagtcata aacaagcaca catttactat gcatacaatg taccgttatg   6463

acaaaggagg accaaaatcc aaacaatatc aaaccacacc aaaaaccaca aggagcctaa   6523

taattactaa ggtgatactt ccaaagggag gactttattt cttagatgag aatgaaaatg   6583

gacacattgg aaattattgg agagccctct ggctatgagt ccttccacaa ccatatggta   6643

ccaccgactg gcaggagaaa tgtgtgaaca tgtgcctcct ctccccaacc actggggccg   6703

gtggggtgac ggtggcactt ttagcagtat cctccgtggt ttgagttgaa aataagtttt   6763

aaaaatcctg tgagtcatgg ttttgcattg aaacctcttc ccactgtgta cccacaaata   6823

gttaactaaa tagaccatta gaaaaggaag aaaatataaa gcagatgcca agcagagatg   6883

tcctaatttt tgacaaaaaa gcaatgttgc ttgtgtcaag aagaaactga actttgtgaa   6943

gagttgaaat ggaattccac tgaattagaa aaacttgttt tctcctgcct ggatacatac   7003
```

8

agtcagggcc attgatgcac aggtgttcct ggctgttgtt acactttacc ctctgaaatg　7063

atgctcccaa gtgctatgtg atgagctcct tgtgtgccca gtggaatagg tgtgtccatg　7123

tgtcatttta aagactatta attacactaa tatagtttct ttctctcttt ggataatag　7182

gca cgt tgt tcc gta tgg act tgg aag ccc tag gtc caa gag agc ctt　7230
Ala Arg Cys Ser Val Trp Thr Trp Lys Pro   Val Gln Glu Ser Leu
        165   170   175

gga gaa ttt act tcc cac aaa ggc aac aga ccg tga gaa tag atg cca　7278
Gly Glu Phe Thr Ser His Lys Gly Asn Arg Pro   Glu   Met Pro
        180   185   190

atg tgc tag cca aaa aga caa gaa gtg ctg gaa ttt ttg cca agc agg　7326
Met Cys   Pro Lys Arg Gln Glu Val Leu Glu Phe Leu Pro Ser Arg
            195   200   205

aaa aga act cag gtgagcagaa acacctttgc ttttcaatca gtttaacagc　7378
Lys Arg Thr Gln
            210

ctcctgaact ccttcctatc atggtactgc cttcctgttt tagagagact aacagagaca　7438

ttgaaagtca gggtaaagct gaatataaca ttgctgaaat gtttttcctt gtgtatttta　7498

acag ggc tga aga cat tat gga gaa aga ctg gaa taa tca taa gaa agg　7547
     Gly   Arg His Tyr Gly Glu Arg Leu Glu   Ser   Glu Arg
                      215   220

aaa aga ctg ttc caa gct tgg gaa aaa gtg tat tta tca gca gtt agt　7595
Lys Arg Leu Phe Gln Ala Trp Glu Lys Val Tyr Leu Ser Ala Val Ser
        225   230   235

gag agg aag aaa aat cag aag aag ttc aga gga aca cct aag aca aac　7643
Glu Arg Lys Lys Asn Gln Lys Lys Phe Arg Gly Thr Pro Lys Thr Asn
    240   245   250

cag gtaagaggga aggaagaaaa attaggtaag aggttcacaa gaacaactag　7696
Gln
255

ccccagtcag tgatgccagc agcctgttcc tccagccctt cttacccggg caggtgaaag　7756

acttagaaaa cagtagcaga ggagatctat gcatcctata gattaaaagg agcaaaagaa　7816

tccctcttaa atatttccat gaagctctgg aatgcaaacc gatgtcctct gtacctttag　7876

cacataccat ttcatctaca ggtagatttc ccaaccaaaa tatatccaga gatgcctttg　7936

tcattgggtt atatacagcc tttgcctctc tgagtcaatg tatttaccac tttccctgag　7996

aaatcgaaaa tcattttggg gagcggacat ttagaaaaag aatcaaagtg tcatggataa　8056

tcaaattctt caataagttg cagttattca gatggccaaa ggaaaaataa agtcattaga　8116

tagggttggt agaatttaga acatgctgtt tttcaggttt atggtctttt tttttttttt   8176

ttttttttt taaataggga aatgtgtttg gtgcagagcc aatgtcattc caaaaagctc   8236

tctcttttcc tggtcagtca tgtgctggga cagagaaggg atctggatta ggcaacatca   8296

tagagttgct ctgagctgct ctttggtgat aacccttcca aatcctaaac ttttttggaat   8356

tcacaagctc aaaggaggaa acctactctc tgatctacca catgttctgc attttttctat   8416

catggtctat ggaaacttct cttagaaatc cagtggcaag aagttctatg attaaagtgt   8476

tctgagctca ggccaggcag tcatgaacta cttctgagtt gtttactact gatttgtggg   8536

gcagcctcag ctatcggttt cttcacacct gcttatgaga gtatccatat ttatggtcgc   8596

aggcagtaat gctccccacg agatcagttt ctgaactaac ctggaatttt ttatgggttt   8656

ttattatgcc aactattaaa tcaacattac agttcttccc tctgtatttc tcctgtaaaa   8716

cattaggcct gcaaaaaaaa aaaatctttt taaaaataat tgccataaag tatttgctct   8776

gggcctactg tatgcttctt ttytttttct ctcttttcaa ctaagtcacc gtcaatttat   8836

taagatggcc ataactattc aaaacctatg ctgagttcct caaggcaggg tcgcatagtg   8896

atgaaggttg ggatggggct acggaagaaa ccagaacaac tctagtttat ttaaaacctg   8956

tatttactgc ccacttcccc ttagacttga ccatatgacc ccttgctccc cattctaagc   9016

atagggcag gctttatttt tacaatggta atagatgata tcacttgagg ttttatcaaa   9076

gagttgcggc gggtggtgaa agttcacaac cagattcagg ttttgtttgt gccagattct   9136

aattttacat gtttcttttg ccaaagggtg atttttttaa aataacattt gttttctctt   9196

atcttgcttt attag gtc gga gac cat gag aaa cag cgt caa atc atc ttt   9247
               Val Gly Asp His Glu Lys Gln Arg Gln Ile Ile Phe
                              260   265

tca tga tcc caa gct gaa agg caa gcc ctc cag aga gcg tta tgt gac   9295
Ser   Ser Gln Ala Glu Arg Gln Ala Leu Gln Arg Ala Leu Cys Asp
            270   275   280

cca caa ccg agc aca ttg gtg aca gac ctt cgg ggc ctg tct gaa gcc   9343
Pro Gln Pro Ser Thr Leu Val Thr Asp Leu Arg Gly Leu Ser Glu Ala
        285   290   295

ata gcc tcc acg gag agc ctg tgg ccg act ctg cac tct cca ccc tgg   9391
Ile Ala Ser Thr Glu Ser Leu Trp Pro Thr Leu His Ser Pro Pro Trp
    300   305   310

ctg gga tca gag cag gag cat cct ctg ctg gtt cct gac tgg caa agg   9439
Leu Gly Ser Glu Gln Glu His Pro Leu Leu Val Pro Asp Trp Gln Arg
315   320   325   330

acc agc gtc ctc gtt caa aac att cca aga aag gtt aag gag ttc ccc   9487

```
Thr Ser Val Leu Val Gln Asn Ile Pro Arg Lys Val Lys Glu Phe Pro
            335                 340                 345


caa cca tct tca ctg gct tcc atc agt ggt aac tgc ttt ggt ctc ttc     9535
Gln Pro Ser Ser Leu Ala Ser Ile Ser Gly Asn Cys Phe Gly Leu Phe
            350                 355                 360


ttt cat ctg ggg atg aca atg gac ctc tca gca gaa aca cac agt cac     9583
Phe His Leu Gly Met Thr Met Asp Leu Ser Ala Glu Thr His Ser His
            365                 370                 375


att cga att cgg gtg gca tcc tcc gga gag aga gag agg aag gag att     9631
Ile Arg Ile Arg Val Ala Ser Ser Gly Glu Arg Glu Arg Lys Glu Ile
            380                 385                 390


cca cac agg ggt gga gtt tct gac gaa ggt cct aag gga gtg ttt gtg     9679
Pro His Arg Gly Gly Val Ser Asp Glu Gly Pro Lys Gly Val Phe Val
395                 400                 405                 410


tct gac tca ggc gcc tgg cac att tca ggg aga aac tcc aaa gtc cac     9727
Ser Asp Ser Gly Ala Trp His Ile Ser Gly Arg Asn Ser Lys Val His
            415                 420                 425


aca aag att ttc taa gga atg cac aaa ttg aaa aca cac tca aaa gac     9775
Thr Lys Ile Phe     Gly Met His Lys Leu Lys Thr His Ser Lys Asp
            430                 435                 440


aaa cat gca agt aaa gaa aaa aaa aag aaa gac ttt tgt tta aat ttg     9823
Lys His Ala Ser Lys Glu Lys Lys Lys Lys Asp Phe Cys Leu Asn Leu
            445                 450                 455


taa aat gca aaa ctg aat gaa act gtt act acc ata aat cag gat atg     9871
    Asn Ala Lys Leu Asn Glu Thr Val Thr Thr Ile Asn Gln Asp Met
            460                 465                 470


ttt cat gaa tat gag tct acc tca cct ata ttg cac tct ggc aga agt     9919
Phe His Glu Tyr Glu Ser Thr Ser Pro Ile Leu His Ser Gly Arg Ser
         ·475                 480                 485


att tcc cac att taa tta ttg cct ccc caa act ctt ccc acc cct gct     9967
Ile Ser His Ile     Leu Leu Pro Pro Gln Thr Leu Pro Thr Pro Ala
    490                 495                 500


gcc cct tcc tcc atc ccc cat act aaa tcc tag cct cgt aga agt ctg    10015
Ala Pro Ser Ser Ile Pro His Thr Lys Ser     Pro Arg Arg Ser Leu
    505                 510                 515


gtc taa tgt gtc agc agt aga tat aat att ttc atg gta atc tac tag    10063
Val     Cys Val Ser Ser Arg Tyr Asn Ile Phe Met Val Ile Tyr
        520                 525                 530


ctc tga tcc ata aga aaa aaa aga tca tta aat cag gag att ccc tgt    10111
Leu     Ser Ile Arg Lys Lys Arg Ser Leu Asn Gln Glu Ile Pro Cys
            535                 540                 545


cct tga ttt ttg gag aca caa tgg tat agg gtt gtt tat gaa ata tat    10159
Pro     Phe Leu Glu Thr Gln Trp Tyr Arg Val Val Tyr Glu Ile Tyr
```

```
                 550                    555                    560

tga aaa gta agt gtt tgt tac gct tta aag cag taa aat tat ttt cct   10207
    Lys Val Ser Val Cys Tyr Ala Leu Lys Gln     Asn Tyr Phe Pro
            565                    570                    575

tta tat aac cgg cta atg aaa gag gtt gga ttg aat ttt gat gta ctt   10255
Leu Tyr Asn Arg Leu Met Lys Glu Val Gly Leu Asn Phe Asp Val Leu
            580                    585                    590

att ttt tta tag ata ttt ata ttc aaa caa ttt att cct tat att tac   10303
Ile Phe Leu     Ile Phe Ile Phe Lys Gln Phe Ile Pro Tyr Ile Tyr
            595                    600                    605

cat gtt aaa tat ctg ttg ggc agg cca tat tgg tct atg tat ttt taa   10351
His Val Lys Tyr Leu Leu Gly Arg Pro Tyr Trp Ser Met Tyr Phe
            610                    615                    620

aat atg tat ttc taa atg aaa ttg aga aca tgc ttt gtt ttg cct gtc   10399
Asn Met Tyr Phe     Met Lys Leu Arg Thr Cys Phe Val Leu Pro Val
            625                    630                    635

aag gta atg act tta gaa aat aaa tat ttt ttt cct tac tgt ac       10443
Lys Val Met Thr Leu Glu Asn Lys Tyr Phe Phe Pro Tyr Cys
            640                    645                    650

tgatttggaa tcattactga aatttgtaag gagtgggcca acgtgattaa gtaccataaa   10503

ggcaaataaa tggttaaaga cggtttcata gaaaagtgac aattagaagg atattacggt   10563

ctaagctaat tatataaaga attttatctg tatcttaaat gttgatttta tactgcattg   10623

aggtaaaaac acaaacaaa aaagcagctt taacacctct gtcttctctt gggtagcagc    10683

ctcctgcttc tccttcacct gaaaaattct ccagggactt catccattaa cttggctcag   10743

gctattggca ggattcacag tttaagctga tggtgtggtg agagatgctt tatccatatt   10803

aatggactga aggaagtaat ggcaagacaa ccccccaaaa catacctaat tatacaaagt   10863

tatataccaa agttgctttt agaaaatggc ctgctcagag caagtagagg tttccaatgg   10923

ctttttattt tctcacatta aggatgttgt ttcttaagga acattgagta ccattgcttc   10983

ttcgtgatag cctaggactg ccgtgtgccc atggaggtag agacaccagg tactgattct   11043

aggtcctctg ccacaaagca ccacttcctc tccactttgc cttggctggc cttgtcagct   11103

cactggagag cacagtattg caattgcagt attgcaaatg gtcactacta actgaattct   11163

ctaagagctt gattagccct cgagaatctt ccttgccctt ctctaatagt gtctgaagga   11223

attcctggca tttaacaaat attagcatgt agtgatcact gtcgtcctaa cagtgacaca   11283

tcagaaggat ttcaaataac agtcttcagg catgcgtaat caatgtcctg tgcagagtct   11343

ccgtcctcat tgatcctcat ttttctcttt aaggcacagt ccaatgtctt tggggaattg   11403
```

```
tttataaagc ttactttatc cataaactgt ttctcagtgc gtgactctga agaaaatttt   11463

gaagttttgc ccatgttgac aaggtgcttg gtctgaactt ggccagtatt taatcttgag   11523

caaacgattc aatttccttc tatcgtgagt tttctcatct atgaaacaag ggagttgagg   11583

ggagtttctt tcatacctct gagaaagagt ttgagattac ataaagaagt tgaagtggca   11643

tgaaaaaaaa taaagatctg agcttagaag acatggatct aatacattta agaggaagtc   11703

agaatcagag aagccactga acaaaacagt ccaaacggag catagtaagt cagattgatg   11763

agttttggtt gggttttttca tcagtcaaac ccttgagccc ccctttccca tgcttcctgc   11823

ttcagtatcc agtaggaaaa atgaaaggga tgatgtagac actctagggc atgaggattt   11883

gcagtaaata agttgggaga ctcacagaaa attaatattt ttcaaacatg aagacgaaac   11943

attcaattat attacagtcc acatcagctt gaagggtaaa ctgatgggat gatctgtcac   12003

atttcttgct ctgtttccag taaaagcatg gtttctggaa acccacttag gacagctttc   12063

tctctttaca ctgatagccc aggcaagctt tgatctcaga actccagaaa ccagagaact   12123

ctaggtggaa tgtggtaact tttgccaggg cagagggaac acctactaat aggtacttca   12183

tttgcaccac cagagattgg catctttttt gatggatcca ctggctttga tactgcctgt   12243

actcccccaa aacacagctt gggtattgga ctaatctaga gctccctcag gagaactctt   12303

gctgacatta agaaagagca acattttgtc tttccaggtg aaaatccaag gccaaaaagg   12363

gagtgactca cctaagatca cagaaggagc tgtagcatct ctggagcctg aacacttaag   12423

ttaagcacga ctatttcacg cagagggcat gaattc                             12459
```

```
<210>   2
<211>   20
<212>   DNA
<213>   Artificial sequence

<220>
<221>   misc_feature
<222>   (1)..(20)
<223>   Primer


<400>   2
ctccatcccc agaaaaactg                                                  20


<210>   3
<211>   20
<212>   DNA
<213>   Artificial sequence
```

```
<220>
<221>  misc_feature
<222>  (1)..(20)
<223>  Primer


<400>  3
aaggaaggtg gtgctgagaa                                                    20


<210>  4
<211>  20
<212>  DNA
<213>  Artificial sequence

<220>
<221>  misc_feature
<222>  (1)..(20)
<223>  Primer


<400>  4
gggggatttc aaggttagat                                                    20


<210>  5
<211>  22
<212>  DNA
<213>  Artificial sequence

<220>
<221>  misc_feature
<222>  (1)..(22)
<223>  Primer


<400>  5
gagaagcccc gataagttct tt                                                 22
```

INTERNATIONAL SEARCH REPORT

International application No.

PCT/US01/28834

A. CLASSIFICATION OF SUBJECT MATTER

IPC(7) : C12Q 1/68; C12P 19/34; C07H 21/02, 21/04

US CL : 435/6, 91.2; 536/23.1, 23.5, 24.31, 24.33

According to International Patent Classification (IPC) or to both national classification and IPC
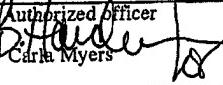
B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

U.S. : 435/6, 91.2; 536/23.1, 23.5, 24.31, 24.33

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)

Please See Continuation Sheet

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X --- Y	STEVENS et al. Genetic Variability of the ET-1 and the ETA Receptor Genes in Essential Hypertension. Journal of Cardiovascular Pharmacology. 1995, Vol. 23, Supplement 3, pages S9-S12, especially pages S11-S12.	1-3, 6, 22, 23, 26 ---------- 7-21
X --- Y	TIRET et al. The Lys198Asn Polymorphism in the Endothelin-1 Gene is Associated with Blood Pressure in Overweight People. Hypertension. 1999, Vol. 33, pages 1169-1174, especially pages 1172-1173.	1-3, 6, 22, 23, 26 ---------- 7-21
Y	US 5,888,819 A (GOELET et al) 30 March 1999, columns 3-5.	7-21

☐ Further documents are listed in the continuation of Box C.   ☐ See patent family annex.

*	Special categories of cited documents:	"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
"A"	document defining the general state of the art which is not considered to be of particular relevance	
"E"	earlier application or patent published on or after the international filing date	"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
"L"	document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)	"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art
"O"	document referring to an oral disclosure, use, exhibition or other means	"&" document member of the same patent family
"P"	document published prior to the international filing date but later than the priority date claimed	

Date of the actual completion of the international search	Date of mailing of the international search report
14 November 2001 (14.11.2001)	2 7 DEC 2001
Name and mailing address of the ISA/US Commissioner of Patents and Trademarks Box PCT Washington, D.C. 20231 Facsimile No. (703)305-3230	Authorized Officer Carla Myers Telephone No. 703-308-0196

Form PCT/ISA/210 (second sheet) (July 1998)

Continuation of B. FIELDS SEARCHED Item 3:
DIALOG: Medline, CA, Biosis, Embase, SciSearch; WEST: US, EP, JP, WO Patents
search terms: EDN-1, endothelin, ET-1, EDN1, preproendothelin, mutation, polymorphism, variant, allele